

AMENDMENTS TO THE SPECIFICATION

Please amend the specification as described below.

Delete paragraphs 0058-0096.

Replace paragraph 0155 with the following paragraph.

The nucleotide sequences of each of a plurality of GAM oligonucleotides that are described by Fig. 1 and their respective genomic sources and genomic locations are set forth in Tables 1-3, hereby incorporated herein. Specifically, in Table 1, line 778 describes GAM RNA (miRNA) as set forth in SEQ ID NO: 348 is shown as predicted from human.

After paragraph 0155, add the following Table 1, paragraph, Table 2, paragraph, and Table 3.

Table 1

GAM SEQ-ID	GAM NAME	GAM RNA SEQUENCE	GAM ORGANISM	GAM POS
=====	=====	=====	=====	=====
348	GAM353678	CAGCAGCACACTGTGGTTTGTA	Human	A

In Table 2, lines 42112-42207, describes GAM PRECURSOR RNA (hairpin) as set forth in SEQ ID NO: 4233864 and as it relates to Figures 1-8 .

Table 2

GAM NAME	GAM ORGA NISM	PRECUR SEQ-ID	PRECURSOR SEQUENCE	GAM DESCRIPTION
=====	=====	=====	=====	=====
GAM 353678	Human	4233 864	CCTGCTCCCG CCCCAGCAGC ACACTGTGGT TTGTACGGCA CTGTGGCCAC GTCCAAACCA CACTGTGGTG TTAGAGCGAG GGTGGGGGAGG	Fig. 1 further provides a conceptual description of another novel bioinformatically-detected human oligonucleotide of the present invention referred to here as the Genomic Address Messenger 353678 (GAM353678) oligonucleotide, which modulates expression of respective target genes whose function and utility are known in the art. GAM353678 is a novel bioinformatically detectable regulatory, non-

protein-coding, miRNA-like oligonucleotide. The method by which GAM353678 is detected is described with additional reference to Figs. 1-8. The GAM353678 precursor, herein designated GAM PRECURSOR, is encoded by the Human genome. GAM353678 target gene, herein designated GAM TARGET GENE, is a target gene encoded by the target organism as specified in Tables 6-7. The GAM353678 precursor, herein designated GAM PRECURSOR, encodes a GAM353678 precursor RNA, herein designated GAM PRECURSOR RNA. Similar to other miRNA oligonucleotides, the GAM353678 precursor RNA does not encode a protein. GAM353678 precursor RNA folds onto itself, forming GAM353678 folded precursor RNA, herein designated GAM FOLDED PRECURSOR RNA, which has a two-dimensional "hairpin" structure. GAM PRECURSOR RNA folds onto itself, forming GAM FOLDED PRECURSOR RNA, which has a two-dimensional "hairpin structure". As is well-known in the art, this "hairpin structure" is typical of RNA encoded by known miRNA precursor oligonucleotides and is due to the full or partial complementarity of the nucleotide sequence of the first half of an miRNA precursor to the RNA that is encoded by a miRNA oligonucleotide to the nucleotide sequence of the second half thereof. A nucleotide sequence that is identical or highly similar to the nucleotide sequence of the GAM353678 precursor RNA is designated SEQ ID NO:4233864, and is provided hereinbelow with reference to the sequence listing section. The nucleotide sequence designated SEQ ID NO:4233864 is located from position 7121806 to position 7121896 relative to chromosome 17 on the "-" strand, and overlaps an intergenic region (UCSC.h16.refGene database). Furthermore, the nucleotide sequence designated SEQ ID NO:4233864 is positioned in a region that is conserved

between human, mouse and rat (UCSC.hgl6.humorMm3Rn3). A schematic representation of a predicted secondary folding of GAM353678 folded precursor RNA, herein designated GAM FOLDED PRECURSOR RNA is set forth in Table 4 incorporated herein. An enzyme complex designated DICER COMPLEX, an enzyme complex composed of Dicer RNaseIII together with other necessary proteins, cuts the GAM353678 folded precursor RNA yielding a single-stranded ~22 nt-long RNA segment designated GAM353678 RNA, herein designated GAM RNA. Table 5 provides a nucleotide sequence that is highly likely to be identical or extremely similar to the nucleotide sequence of GAM353678 RNA, hereby incorporated herein. GAM353678 target gene, herein designated GAM TARGET GENE, encodes a corresponding messenger RNA, designated GAM353678 target RNA, herein designated GAM TARGET RNA. As is typical of mRNA of a protein-coding gene, GAM353678 target RNA comprises three regions, as is typical of mRNA of a protein-coding gene: a 5' untranslated region, a protein-coding region and a 3' untranslated region, designated 5'UTR, PROTEIN-CODING and 3'UTR, respectively. GAM353678 RNA, herein designated GAM RNA, binds complementarily to one or more target binding sites located in the untranslated regions of GAM353678 target RNA. This complementary binding is due to the partial or full complementarity between the nucleotide sequence of GAM353678 RNA and the nucleotide sequence of each of the target binding sites. As an illustration, Fig. 1 shows three such target binding sites, designated BINDING SITE I, BINDING SITE II and BINDING SITE III, respectively. It is appreciated that the number of target binding sites shown in Fig. 1 is only illustrative and that any suitable number of target binding sites may be present. It is further appreciated that although Fig.

1 shows target binding sites only in the 3'UTR region, these target binding sites may instead be located in the 5'UTR region or in both the 3'UTR and 5'UTR regions. The complementary binding of GAM353678 RNA, herein designated GAM RNA, to target binding sites on GAM353678 target RNA, herein designated GAM TARGET RNA, such as BINDING SITE I, BINDING SITE II and BINDING SITE III, inhibits the translation of GAM353678 target RNA into respective GAM353678 target protein, herein designated GAM TARGET PROTEIN, shown surrounded by a broken line. It is appreciated that the GAM353678 target gene, herein designated GAM TARGET GENE, in fact represents a plurality of GAM353678 target genes. The mRNA of each one of this plurality of GAM353678 target genes comprises one or more target binding sites, each having a nucleotide sequence which is at least partly complementary to GAM353678 RNA, herein designated GAM RNA, and which when bound by GAM353678 RNA causes inhibition of translation of the GAM353678 target mRNA into a corresponding GAM353678 target protein. The mechanism of the translational inhibition that is exerted by GAM353678 RNA, herein designated GAM RNA, on one or more GAM353678 target genes, herein collectively designated GAM TARGET GENE, may be similar or identical to the known mechanism of translational inhibition exerted by known miRNA oligonucleotides. The nucleotide sequence of GAM353678 precursor RNA, herein designated GAM PRECURSOR RNA, its respective genomic source and genomic location and a schematic representation of a predicted secondary folding of GAM353678 folded precursor RNA, herein designated GAM FOLDED PRECURSOR RNA are set forth in Tables 3-4, hereby incorporated herein. The nucleotide sequences of a "diced" GAM353678 RNA, herein

designated GAM RNA, from GAM353678 folded precursor RNA are set forth in Table 5, hereby incorporated herein. The nucleotide sequences of target binding sites, such as BINDING SITE I, BINDING SITE II and BINDING SITE III of Fig. 1, found on GAM353678 target RNA, herein designated GAM TARGET RNA, and a schematic representation of the complementarity of each of these target binding sites to GAM353678 RNA, herein designated GAM RNA, are set forth in Tables 6-7, hereby incorporated herein. It is appreciated that the specific functions and accordingly the utilities of GAM353678 RNA are correlated with and may be deduced from the identity of the GAM353678 target gene inhibited thereby, and whose functions are set forth in Table 8, hereby incorporated herein.

Table 3, lines 1279-1280, shows data relating to the source and location of the GAM oligonucleotide, specifically the GAM PRECURSOR (hairpin) and its position in the genomic sequence of human.

Table 3

GAM NAME	PRECUR SOR SEQ-ID	GAM ORGANISM	SOURCE	STR AND	SRC-START OFFSET	SRC-END OFFSET
=====	=====	=====	=====	=====	=====	=====
GAM353678	4233864	Human	17		7121806	7121896

Replace paragraph 0156 with the following paragraph.

The nucleotide sequences of GAM PRECURSOR RNAs, and a schematic representation of a predicted secondary folding of GAM FOLDED PRECURSOR RNAs, of each of a plurality of GAM oligonucleotides described by Fig. 1 are set forth in Table 4, hereby incorporated herein. Table 4 lines 2384-2388, shows a schematic representation of the GAM folder precursor as set forth in SEQ ID NO:348, beginning at the 5' end (beginning of upper row) to the 3'

end (beginning of lower row), where the hairpin loop is positioned at the right part of the drawing.

After paragraph 0156, add the following Table 4.

Table 4

GAM NAME	PRE CUR SEQ -ID	GAM ORGA NISM	PRECURSOR SEQUENCE	GAM FOLDED PRECURSOR RNA									
====	====	=====	=====	=====									
GAM 353 678	423 386 4	Human	CCTGCTCCCGCCCCAGCAGC ACACTGTGGTTTGTACGGCA CTGTGGCCACGTCCAAACCA CACTGTGGTGTAGAGCGAG GGTGGGGGAGG	G	C-----	G	C	T	--	AC			
				CCT	CTCCCGCCC	AGCA	CACA	TGTGGTTTG	AC	GGC	T		
				GGA	GGGGGTGGG	TTGT	GTGT	ACACCAAAC	TG	CCG	G		
				-	AGCGAGA	G	C		C	CA	GT		

Replace paragraph 0157 with the following paragraph.

The nucleotide sequences of “diced” GAM RNAs of each of a plurality of GAM oligonucleotides described by Fig. 1 are set forth in Table 5, hereby incorporated herein. Table 5, line 1276 shows the mature GAM RNA as set forth in SEQ ID NO: 348 as sliced by DICER from the GAM PRECURSOR sequence (hairpin) as set forth in SEQ ID NO: 4233864.

After paragraph 0157, add the following Table 5.

GAM NAME	GAM ORGANISM	GAM RNA SEQUENCE	PRECUR SEQ-ID	GAM POS
=====	=====	=====	=====	=====
GAM353678	Human	CAGCAGCACACTGTGGTTTGT	4233864	A

Replace paragraph 0158 with the following paragraph.

The nucleotide sequences of target binding sites, such as BINDING SITE I, BINDING SITE II and BINDING SITE III that are found on GAM TARGET RNAs of each of a plurality of GAM oligonucleotides that are described by Fig. 1, and a schematic representation of the complementarity of each of these Target binding sites to each of a plurality of GAM RNAs that are described by Fig. 1 are set forth in Tables 6-7, hereby incorporated herein. Table 6 shows data relating to the SEQ ID NO of the GAM target binding site sequence of the target gene name as bound by the GAM RNA as set forth in SEQ ID NO: 348. Table 6, lines 3688165, 767082, 762322 and 763042 related to target binding site SEQ ID NO: 1810388, 673420, 671402 respectively.

After paragraph 0158, add the following Table 6, paragraph, and Table 7.

Table 6

TARGET BINDING SITE	TARGET SEQ-ID	TARGET ORGANISM	TARGET	TARGET BINDING SITE SEQUENCE
=====	=====	=====	=====	=====
1810388		Human	MGAT5	CACCATGCTGCTG
673420		Human	SERPINH1	AACTAGGTGCTGCAG
671402		Human	SERPINH1	ATACCATGATGCTG
671042		Human	SERPINH1	CTATAAACTAGGTGCTGCAG

Table 7, lines 312839-313773 shows data relating to target genes and binding site of GAM oligonucleotides.

Table 7

GAM NAME	GAM ORGA NISM	GAM RNA SEQUENCE	TARGET BS-SEQ	TARG ET	TARGET REF-ID	TARGET ORGANISM	UTR	BINDING SITE (UPPER:TARGET; LOWER:GAM)	DRAW	GAM POS
=====	=====	=====	=====	=====	=====	=====	=====	=====	=====	=====
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	AAACCAAA CTTATGCA GCTG	nup C	NC_004431 f rom 27953 90 to 27966 31 (+)	Escher ichia coli CFT073	3 ---	A C TA A AAACCA A T TGC GCTG TTTGGT T A ACG CGAC ATG G C C- A		A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	AAACCAAA CTTATGCA GCTG	nup C	NC_004741 f rom 24940 19 to 24952 21 (+)	Shigell a file xneri 2a str . 2457T	3 ---	A C TA A AAACCA A T TGC GCTG TTTGGT T A ACG CGAC ATG G C C- A		A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	AAACCTTG CTGCG	rel A	NC_000962 f rom 29078 24 to 29101 96 (-)	Mycobac teriu m tubercu los is H37Rv	3 ---	C----- AAACC TGCTGC G TTTGG ACGACG C ATG TGTCAC A		A

GAM35	Hum	CAGCAGCA	AAACCCCTG	rel	NC_002945 f	Mycobac	3	---	C-----	-	A
3678	an	CACGTGTGG	CTGCG	A	rom 28752	teriu m		AAACC	TGCTGC	G	
		TTTGTA			74 to 28776	bovis		TTTGG	ACGACG	C	
					46 (-)	subs p		ATG	TGTCAC	A	
						bovis					
						AF21					
						22/97					
GAM35	Hum	CAGCAGCA	AAACCCCTT	yab	NC_004431 f	Escheri	3	---	C TTC-	T	A
3678	an	CACGTGTGG	TCTGTCTGC	O	rom 614	chia		AAACC T	TGCTGCT		
		TTTGTA	TT		89 to 62148	coli		TTTGG G	ACGACGA		
					(-)	CFT073		ATG	T TCAC	C	
GAM35	Hum	CAGCAGCA	AAACCGAT	amt	NC_004337 f	Shigell	3	---	AT CAG	G	A
3678	an	CACGTGTGG	GCAGTGC	B	rom 4080	a fle		AAACCG G	TGC GCTG		
		TTTGTA	GCTG		59 to 40934	xneri		TTTGGT C	ACG CGAC		
					5 (+)	2a str		ATG	GT AC-	A	
						. 301					
GAM35	Hum	CAGCAGCA	AAACCGAT	amt	NC_004741 f	Shigell	3	---	AT CAG	G	A
3678	an	CACGTGTGG	GCAGTGC	B	rom 4078	xneri		AAACCG G	TGC GCTG		
		TTTGTA	GCTG		60 to 40914	2a str		TTTGGT C	ACG CGAC		
					6 (+)	. 2457T		ATG	GT AC-	A	
GAM35	Hum	CAGCAGCA	AAACCGAT	amt	NC_004431 f	Escheri	3	---	AT CCG	G	A
3678	an	CACGTGTGG	GCCGTGCG	B	rom 5476	chia		AAACCG G	TGC GCTG		
		TTTGTA	GCTG		16 to 54890	coli		TTTGGT C	ACG CGAC		
					2 (+)	CFT073		ATG	GT AC-	A	
GAM35	Hum	CAGCAGCA	AAACCGCC	dsd	NC_003197 f	Salmone	3	---	CCCCAG	-	A
3678	an	CACGTGTGG	CCCAGTCT	A	rom 40044	lla t		AAACCGC	T		
		TTTGTA	GCTG		53 to 40057	yphimur		CTGCTG	TTTGGTG		
					75 (+)	ium L		A	GACGAC	ATG	
						T2		TCAC--	C		
GAM35	Hum	CAGCAGCA	AAACCGGC	gad	NC_002947 f	Pseudom	3	---	GCC -	C	A
3678	an	CACGTGTGG	CTTGCCGC		rom 48716	onas		AAACCG	T TGC	GCTG	
		TTTGTA	TG		25 to 48729	putida		TTTGGT	A ACG	CGAC	
					29 (+)	KT244 0		ATG	GTC C	A	
GAM35	Hum	CAGCAGCA	AAACGAAT	aro	NC_003116 f	Neisser	3	---	G ATTGAATCA	C	A
3678	an	CACGTGTGG	TGAATCAT	A	rom 15575	ia me		AAAC A	TGC		
		TTTGTA	GCCGCTG		02 to 15588	ningiti		GCTG	TTTG T		
					03 (+)	dis Z		ACG	CGAC	ATG	
						2491		G	GTCAC----	A	
GAM35	Hum	CAGCAGCA	AAACGACG	ruv	NC_003143 f	Yersini	3	---	G ---G	AAAC	A
3678	an	CACGTGTGG	GGCTGCTG	B	rom 23364	a pes		ACG	GCTGCTG	TTTG	
		TTTGTA			49 to 23374	tis		TGT	CGACGAC	ATG	G
					53 (+)			CACA			
GAM35	Hum	CAGCAGCA	AAACGACG	ruv	NC_004088 f	Yersini	3	---	G ---G	AAAC	A
3678	an	CACGTGTGG	GGCTGCTG	B	rom 24820	a pes		ACG	GCTGCTG	TTTG	
		TTTGTA			31 to 24830	tis KIM		TGT	CGACGAC	ATG	G
					35 (-)			CACA			
GAM35	Hum	CAGCAGCA	AAACGATA	yci	NC_004431 f	Escheri	3	---	G --- C		A
3678	an	CACGTGTGG	TCCTGCTG	E	rom 15586	chia		AAAC ATA	T CTGCTG		
		TTTGTA			41 to 15591	coli		TTTG TGT	A GACGAC		
					47 (-)	CFT073		ATG	G CAC C		
GAM35	Hum	CAGCAGCA	AAACGCAT	van	NC_002516 f	Pseudom	3	---	G T CA	-	A
3678	an	CACGTGTGG	GTTCATGC	B	rom 55041	onas		AAAC	CATG T	TGC	
		TTTGTA	GCTG		20 to 55050	aerugin		GCTG	TTTG GTGT	A	
					73 (+)	osa P		ACG	CGAC	ATG	C
						A01		C-	A		
GAM35	Hum	CAGCAGCA	AAACGCTC	fts	NC_000922 f	Chlamyd	3	---	GC CCGTATAC		A
3678	an	CACGTGTGG	CGTATACT	Y	rom 11154	ophil a		A	AAAC	T	

	TTTGTA	GCTGCTA		27 to 11162	pneumon		TGCTGCT	TTTG
				99 (-)	iae	G	ACGACGA	ATG
					CWL029		GT TCAC----	C
GAM35	Hum	CAGCAGCA	fts	NC_002491 f	Chlamyd	3	GC CCGTATAC	
3678	an	CACGTGTT	Y	rom 11131	ophil a		A AAAC T	A
		TTTGTA		27 to 11139	pneumon		TGCTGCT	TTTG
				99 (-)	iae	G	ACGACGA	ATG
					J138		GT TCAC----	C
GAM35	Hum	CAGCAGCA	rbs	NC_004337 f	Shigell	3	ATCGACAGT	
3678	an	CACGTGTT	R	rom 39477	a file		- AAAC T	A
		TTTGTA		08 to 39487	xneri		TGCTGC G	
				00 (+)	2a str		TTTGGT	
					. 301		ACGACG C	ATG
							GTCAC----	A
GAM35	Hum	CAGCAGCA	rbs	NC_004741 f	Shigell	3	ATCGACAGT	
3678	an	CACGTGTT	R	rom 38245	a file		- AAAC T	A
		TTTGTA		94 to 38255	xneri		TGCTGC G	
				77 (-)	2a str		TTTGGT	
					. 2457T		ACGACG C	ATG
							GTCAC----	A
GAM35	Hum	CAGCAGCA	SER			3	G ---	A
3678	an	CACGTGTT	PIN	NM_001235	Human		AAAC T G	TGCTGC G
		TTTGTA	H 1				TTTGGT T	ACGACG C
							ATG G CAC	A
GAM35	Hum	CAGCAGCA	aro	NC_004337 f	Shigell	3	T G C GCAA	
3678	an	CACGTGTT	H	rom 15575	a file		AAAC CA G TG	A
		TTTGTA		27 to 15585	xneri		GCTGCTG	TTTG GT T
				73 (-)	2a str		AC CGACGAC	ATG -
					. 301		G C ---A	
GAM35	Hum	CAGCAGCA	zra	NC_003197 f	Salmone	3	G -----	T
3678	an	CACGTGTT	P	rom 43877	lla t		AAA C	TGCTGCT
		TTTGTA		27 to 43881	yphimur		TTT G	ACGACGA
				82 (-)	ium L		ATG G	TGTCAC
					T2			C
GAM35	Hum	CAGCAGCA	cys	NC_003197 f	Salmone	3	GTGTACT T	
3678	an	CACGTGTT	M	rom 25516	lla t		AAATCA	TG
		TTTGTA		51 to 25525	yphimur		TGCTG	TTTGGT
				62 (-)	ium L		AC ACGAC	ATG
					T2		GTCAC---	G
GAM35	Hum	CAGCAGCA	rbs	NC_004431 f	Escheri	3	- - - - -	AA C
3678	an	CACGTGTT	R	rom 44392	chia		AT	TGCTGCTG
		TTTGTA		60 to 44402	coli		TG	ACGACGAC
				52 (+)	CFT073		TCAC	ATG T G
GAM35	Hum	CAGCAGCA	oxy	NC_003197 f	Salmone	3	- - - - -	C
3678	an	CACGTGTT	R	rom 43430	lla t		AA C	TGCTGCT
		TTTGTA		80 to 43439	yphimur		TT G	ACGACGA
				97 (+)	ium L		ATG T	GIGTCAC
					T2			C
GAM35	Hum	CAGCAGCA	oxy	NC_003198 f	Salmone	3	- - - - -	C
3678	an	CACGTGTT	R	rom 36072	lla e		AA C	TGCTGCT
		TTTGTA		04 to 36081	nterica		TT G	ACGACGA
				21 (-)	rica		ATG T	GIGTCAC
					serovar			C
					Typhi			
GAM35	Hum	CAGCAGCA	oxy	NC_004631 f	Salmone	3	- - - - -	C
3678	an	CACGTGTT	R	rom 35928	lla e		AA C	TGCTGCT
		TTTGTA		64 to 35937	nterica		TT G	ACGACGA
				81 (-)	rica		ATG T	GIGTCAC
					serovar			C

				Typhi Ty2					
GAM35	Hum	CAGCAGCA	AAGCCGGT	aro	NC_003116 f	Neisser	---	GTTGCGG	
3678	an	CACTGTGG	TGCGGTGC	A	rom 15575	ia me 3	AAGCCG		A
		TTTGTA	TGCTG		02 to 15588	ningiti	TGCTGCTG		
					03 (+)	dis Z	TTTGGT		
						2491	ACGACGAC ATG		
							GTCAC--		
				Salmone					
GAM35	Hum	CAGCAGCA	AATCCACT	glg	NC_003198 f	lla e 3	---	T TCC T AA	A
3678	an	CACTGTGG	CCGTGTTG	P	rom 41445	nterica	CCAC	GTG TGCTG TT	
		TTTGTA	CTG		68 to 41470	ente	GGTG	CAC ACGAC ATG T	
					15 (+)	rica	TCA G		
						serovar			
						Typhi			
				Salmone					
GAM35	Hum	CAGCAGCA	AATCCACT	glg	NC_004631 f	lla e 3	---	T TCC T AA	A
3678	an	CACTGTGG	CCGTGTTG	P	rom 41292	nterica	CCAC	GTG TGCTG TT	
		TTTGTA	CTG		15 to 41316	ente	GGTG	CAC ACGAC ATG T	
					62 (+)	rica	TCA G		
						serovar			
						Typhi			
				Ty2					
GAM35	Hum	CAGCAGCA	ACATGCTG	nup	NC_004431 f	Escheri 3	---	- - - - -	T
3678	an	CACTGTGG	CTT	C	rom 27953	chia	A C A	TGCTGCT	A
		TTTGTA			90 to 27966	coli	T G T	ACGACGA	
					31 (+)	CFT073	ATG TT G GTCAC		C
				Shigell					
GAM35	Hum	CAGCAGCA	ACATGCTG	nup	NC_004741 f	a file 3	---	- - - - -	T
3678	an	CACTGTGG	CTT	C	rom 24940	xneri	A C A	TGCTGCT	A
		TTTGTA			19 to 24952	2a str	T G T	ACGACGA	
					21 (+)	. 2457T	ATG TT G GTCAC		C
				Mycobac					
GAM35	Hum	CAGCAGCA	ACGATGGT	pho	NC_000962 f	teriu m 3	---	- - G TAC	
3678	an	CACTGTGG	GTACTGCT	Y2	rom 9135	teriu m	T A C	ATGGTG A	
		TTTGTA	GCTT		56 to 91419	tubercu	TGCTGCT T G		
					7 (-)	los is	TGTCAC ACGACGA ATG TT		
						H37Rv	G --- C		
				Mycobac					
GAM35	Hum	CAGCAGCA	ACGATGGT	pho	NC_002945 f	teriu m 3	---	- - G TAC	
3678	an	CACTGTGG	GTACTGCT	Y2	rom 9143	bovis	T A C	ATGGTG A	
		TTTGTA	GCTT		88 to 91502	subs p	TGCTGCT T G		
					9 (-)	bovis	TGTCAC ACGACGA ATG TT		
						AF21	G --- C		
						22/97			
				Salmone					
GAM35	Hum	CAGCAGCA	ACTGCTGC	glg	NC_003198 f	lla e 3	---	- - - - -	C
3678	an	CACTGTGG	TC	P	rom 41445	nterica	A C	TGCTGCT	A
		TTTGTA			68 to 41470	ente	T G	ACGACGA	
					15 (+)	rica	ATG TT GTGTCAC		C
						serovar			
						Typhi			
				Salmone					
GAM35	Hum	CAGCAGCA	ACTGCTGC	glg	NC_004631 f	lla e 3	---	- - - - -	C
3678	an	CACTGTGG	TC	P	rom 41292	nterica	A C	TGCTGCT	A
		TTTGTA			15 to 41316	ente	T G	ACGACGA	
					62 (+)	rica	ATG TT GTGTCAC		C
						serovar			
						Typhi			
				Ty2					
GAM35	Hum	CAGCAGCA	AGAAITGT	rec	NC_002677 f	Mycobac 3	---	- - T TTAG	A
3678	an	CACTGTGG	GTTAGTGC	G	rom 20147	teriu m	AGA	ATG TG	

	TTTGTA	TGCTG		23 to 20169 54 (-)	leprae		TGCTGCTG	TTT	TGT	
							AC	ACGACGAC	ATG	GG
							C	----		
GAM35	Hum	CAGCAGCA		NC_000962 f	Mycobac	3	---	AA	AACTG	
3678	an	CACTGTGG	rel	rom 29078	teriu m			AGACCATG	G	A
		TTTGTA	A	24 to 29101	tubercu			GCTGCTG	TTTGGTGT	
				96 (-)	los is			C	CGACGAC	ATG
					H37Rv				CA	----A
					Mycobac					
GAM35	Hum	CAGCAGCA		NC_002945 f	teriu m	3	---	AA	AACTG	
3678	an	CACTGTGG	rel	rom 28752	bovis			AGACCATG	G	A
		TTTGTA	A	74 to 28776	subs p			GCTGCTG	TTTGGTGT	
				46 (-)	bovis			C	CGACGAC	ATG
					AF21				CA	----A
					22/97					
GAM35	Hum	CAGCAGCA		NC_003143 f	Yersini	3	---	-	-----	AG CT
3678	an	CACTGTGG	glp	rom 42896	a pes			TGCTGCTG	TT	GG
		TTTGTA	C	50 to 42908	tis			ACGACGAC	ATG	T
				97 (-)				TGTCAC		
GAM35	Hum	CAGCAGCA		NC_004088 f	Yersini	3	---	-	-----	AG CT
3678	an	CACTGTGG	glp	rom 4546	a pes			TGCTGCTG	TT	GG
		TTTGTA	C	77 to 45604	tis KIM			ACGACGAC	ATG	T
				7 (+)				TGTCAC		
					Chlamyd					
GAM35	Hum	CAGCAGCA		NC_000922 f	ophil a	3	---	T	A----	G
3678	an	CACTGTGG	fts	rom 11154	pneumon			ACCA	GCTGCTG	T
		TTTGTA	Y	27 to 11162	iae			TGGT	CGACGAC	ATG
				99 (-)	CWL029				GTCACA	
					Chlamyd					
GAM35	Hum	CAGCAGCA		NC_002491 f	ophil a	3	---	T	A----	G
3678	an	CACTGTGG	fts	rom 11131	pneumon			ACCA	GCTGCTG	T
		TTTGTA	Y	27 to 11139	iae			TGGT	CGACGAC	ATG
				99 (-)	J138				GTCACA	
					Human					
GAM35	Hum	CAGCAGCA		SER		3	---	T	-----	A
3678	an	CACTGTGG	PIN	NM_001235				ACCA	TG	TGCTG
		TTTGTA	H 1					TGGT	AC	ACGAC
									ATG	T
								GTCAC	G	
					Leptosp					
GAM35	Hum	CAGCAGCA		NC_004342 f	ira i	3	---	TT	-----	T
3678	an	CACTGTGG	aro	rom 481	nterrog			A	TC	TGCTGCT
		TTTGTA	D	28 to 48832	ans s			T	GG	ACGACGA
				(-)	erovar			ATG	TT	TGTCAC
					lai s					C
					tr.					
					56601					
GAM35	Hum	CAGCAGCA		NC_002929 f	Bordete	3	--	A-	C	C
3678	an	CACTGTGG	acc	rom 9264	lla p			CAAA	ACGGT	TGC
		TTTGTA	C	07 to 92777	ertussi			GTTT	TGTC	ACG
				7 (+)	s			AT	GG	C
										A
GAM35	Hum	CAGCAGCA		NC_003197 f	Salmone	3	--	--	GCT	TTGCT
3678	an	CACTGTGG	zra	rom 43877	lla t			CAAA	A	T
		TTTGTA	P	27 to 43881	yphimur			GCTG	GTTT	T
				82 (-)	ium L				ACG	CGAC
					T2			GTC	C-----	A
										G
GAM35	Hum	CAGCAGCA		NC_004310 f	Bruce	3	--	--	GTCGTCCTG	
3678	an	CACTGTGG	glp	rom 2107	a sui			CAAA	A	TGC
		TTTGTA	D	63 to 21227	s			GCTG	GTTT	T
				4 (+)	1330				ACG	CGAC
									AT	
									GG	GTCAC-----
										A
GAM35	Hum	CAGCAGCA		NC_004337 f	Shigell	3	--	-	G	C--
3678	an	CACTGTGG	avt	rom 37211	a file			A	G	TGCTGCTG
		TTTGTA	G A	75 to 37225	xneri			T	T	ACGACGAC
										AT
										G

			33	(+)	2a str	G CAC	
					. 301		
					Shigell		
GAM35	Hum	CAGCAGCA	CAAACAGG	avt	NC_004741 f	a file 3	-- - G C-- CAAAC A
3678	an	CACGTGTG	CTGCTGCT	G A	rom 40526	xneri	A G TGCTGCTG GTTTG
		TTTGTA			85 to 40539	2a str	T T ACGACGAC AT G
					38 (-)	. 2457T	G CAC
GAM35	Hum	CAGCAGCA	CAAACATC	mia	NC_000117 f	Chlamyd 3	-- - CA GT T
3678	an	CACGTGTG	ATGGTTC	A	rom 8992	ia tr	CAAAC AT TG TGCTG A
		TTTGTA	TGTTG		76 to 90029	achomat	TG GTTTG TG AC
					5 (+)	is	ACGAC AC AT G TC
							-- G
GAM35	Hum	CAGCAGCA	CAAACCAG	sel	NC_002947 f	Pseudom 3	-- GC G- -
3678	an	CACGTGTG	CGGTCTGC	B	rom 5821	onas	CAAACCA G T CTGCTG A
		TTTGTA	TG		33 to 58405	putida	GTITGGT C A GACGAC
					5 (+)	KT244 0	AT GT AC C
GAM35	Hum	CAGCAGCA	CAAACCAT	mia	NC_000117 f	Chlamyd 3	-- ----- A
3678	an	CACGTGTG	GAIGCTG	A	rom 8992	ia tr	CAAACCA TG TGCTG A
		TTTGTA			76 to 90029	achomat	GTITGGT AC ACGAC
					5 (+)	is	AT GTCAC G
GAM35	Hum	CAGCAGCA	CAAACCGA	min	NC_002947 f	Pseudom 3	-- ACCC-
3678	an	CACGTGTG	CCCTGCTG	E	rom 19326	onas	CAAACCG TGCTGCTG A
		TTTGTA	CTG		80 to 19329	putida	GTITGGT ACGACGAC
					34 (-)	KT244 0	AT GTCAC
GAM35	Hum	CAGCAGCA	CAAACCGC	dna	NC_002677 f	Mycobac 3	-- AC G
3678	an	CACGTGTG	AGTACTGG	E	rom 14230	teriu m	CAAACCGCAGT TG TGCTG A
		TTTGTA	TGCTG		14 to 14265	leprae	GTITGGTGTCA AC ACGAC
					47 (+)		AT C- G
						Leptosp	
						ira i	
GAM35	Hum	CAGCAGCA	CAAACCTCT	aro	NC_004342 f	nterrog 3	-- C TT TC T
3678	an	CACGTGTG	TTTTCTTC	D	rom 481	ans s	CAAACCT T T T CTGCTG A
		TTTGTA	TGCTG		28 to 48832	erovar	GTITGG G A A GACGAC
					(-)	lai s	AT T TC C- C
						tr.	
						56601	
						Salmone	
GAM35	Hum	CAGCAGCA	CAAAGCAC	pts	NC_003198 f	lla e 3	-- G ---- CAAA A
3678	an	CACGTGTG	TGCTGCTG	H	rom 25054	nterica	CAC TGCTGCTG GTTT
		TTTGTA			03 to 25056	ente	GTG ACGACGAC AT G
					60 (+)	rica	TCAC
						serovar	
						Typhi	
GAM35	Hum	CAGCAGCA	CAAAGCCG	amt	NC_004337 f	Shigell 3	-- G CGC -- -
3678	an	CACGTGTG	CGTGCCT	G B	rom 4080	a file	CAAA C G TGC GCTG A
		TTTGTA			59 to 40934	xneri	GTIT G C ACG CGAC
					5 (+)	2a str	AT G TGT AC A
						. 301	
GAM35	Hum	CAGCAGCA	CAAAGCCG	amt	NC_004431 f	Escheri 3	-- G CGC -- -
3678	an	CACGTGTG	CGTGCCT	G B	rom 5476	chia	CAAA C G TGC GCTG A
		TTTGTA			16 to 54890	coli	GTIT G C ACG CGAC
					2 (+)	CFT073	AT G TGT AC A
						Shigell	
GAM35	Hum	CAGCAGCA	CAAAGCCG	amt	NC_004741 f	a file 3	-- G CGC -- -
3678	an	CACGTGTG	CGTGCCT	G B	rom 4078	xneri	CAAA C G TGC GCTG A
		TTTGTA			60 to 40914	2a str	GTIT G C ACG CGAC
					6 (+)	. 2457T	AT G TGT AC A
GAM35	Hum	CAGCAGCA	CAAAGCCT	pil	NC_002947 f	Pseudom 3	-- G C TT TTCGG A

3678	an	CACGTGG TTTGTA	TTTTTCGG GCTGCTG	T	rom	58169 34 to 58179 44 (-)	onas putida KT244 0	CAAA C T T GCTGCTG GTTT G G A CGACGAC AT G T TC C---A
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CAAAGCGT CATGTAAT GCTTGCTG	cys M	NC_003197 f rom 25516 51 to 25525 62 (-)	Salmone lla t yphimur ium L T2	3 -- G CA TAA T CAAA CGT TG TGCT A GCTG GTTT GTG AC ACGA CGAC AT G TC --- -	
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CAAATCCC CAGTTGTG CTG	glc C	NC_004431 f rom 35428 71 to 35436 95 (+)	Escheri chia coli CFT073	3 -- C CAGT - CAAATC C TG TGCTG A GTTTGG G AC ACGAC AT T TCAC G	
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CAACACAT TACTGCTT GCTCTG	aro H	NC_004337 f rom 15575 27 to 15585 73 (-)	Shigell a file xneri 2a str . 301	3 -- CA TACTGCT - CAA CAT TGCT A CTG GTT GTG ACGA GAC AT TG TCAC--- C	
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CAACACTT TGCGCTG	spe D	NC_003197 f rom 1942 01 to 19499 5 (-)	Salmone lla t yphimur ium L T2	3 -- - - TT-- - CAA C AC TGC GCTG A GTT G TG ACG CGAC AT T G TCAC A	
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CAACACTT TGCGCTG	spe D	NC_003198 f rom 1963 89 to 19718 3 (-)	Salmone lla e nterica ente rica serovar Typhi	3 -- - - TT-- - CAA C AC TGC GCTG A GTT G TG ACG CGAC AT T G TCAC A	
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CAACACTT TGCGCTG	spe D	NC_004631 f rom 1963 80 to 19717 4 (-)	Salmone lla e nterica ente rica serovar Typhi Ty2	3 -- - - TT-- - CAA C AC TGC GCTG A GTT G TG ACG CGAC AT T G TCAC A	
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CAACAGCA GTTGCTGC TG	ace K	NC_002947 f rom 51847 42 to 51864 57 (-)	Pseudom onas putida KT244 0	3 -- - - GCAGT CAA C A TGCTGCTG GTT G A T ACGACGAC AT T G GTCAC	
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CAACAGTT TCTTGGTC TGCTG	fmt B(m r p)	NC_002745 f rom 22181 45 to 22255 90 (-)	Staphyl ococc us aureus su bsp. aureus N315	3 -- - - GTT CTTGG - CAA C A T T A CTGCTG GTT G T A A GACGAC AT T G GTC C---- C	
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CAACAGTT TCTTGGTC TGCTG	fmt B(m r p)	NC_002758 f rom 22879 35 to 22953 80 (-)	Staphyl ococc us aureus su bsp. aureus Mu50	3 -- - - GTT CTTGG - CAA C A T T A CTGCTG GTT G T A A GACGAC AT T G GTC C---- C	
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CAACAGTT TCTTGGTC TGCTG	tru nca t ed fmt	NC_003923 f rom 22380 83 to 22401 43 (-)	Staphyl ococc us aureus su bsp.	3 -- - - GTT CTTGG - CAA C A T T A CTGCTG GTT G T A A GACGAC AT T G GTC C---- C	

			B	aureus MW2	
GAM35	Hum	CAGCAGCA	CAACCCAC	cys NC_002947 f	Pseudom 3 -- C CAGCAC -
3678	an	CACTGTGG	CAGCACTG	rom 3151	CAA CCAC TGCTGC A
		TTTGTA	CTGCG	84 to 31598	G GTT GGTT
				4 (+)	ACGACG C AT T
					TCAC-- A
GAM35	Hum	CAGCAGCA	CAACCGCT	pho NC_000962 f	Mycobac 3 -- - ---- CAA
3678	an	CACTGTGG	GCTGCTG	rom 9135	CCGC TGCTGCTG GTT A
		TTTGTA		56 to 91419	GGTG ACGACGAC AT T
				7 (-)	TCAC
GAM35	Hum	CAGCAGCA	CAACCGCT	pho NC_002945 f	Mycobac 3 -- - ---- CAA
3678	an	CACTGTGG	GCTGCTG	rom 9143	CCGC TGCTGCTG GTT A
		TTTGTA		88 to 91502	GGTG ACGACGAC AT T
				9 (-)	TCAC
GAM35	Hum	CAGCAGCA	CAACCGGT	dad NC_004431 f	Escheri 3 -- - G---- -
3678	an	CACTGTGG	GCTGCG	rom 14763	CAA CCG TGCTGC G A
		TTTGTA		06 to 14773	GTT GGT ACGACG C
				76 (+)	AT T GTCAC A
GAM35	Hum	CAGCAGCA	CAACCGTC	fha NC_002929 f	Bordete 3 -- - C GTGA -
3678	an	CACTGTGG	GGTGATGC	rom 30858	CAA CCGT G TGCT A
		TTTGTA	TCTG	65 to 30984	CTG GTT GGTTG C
				55 (+)	ACGA GAC AT T T
					AC-- C
GAM35	Hum	CAGCAGCA	CAACCTGC	spe NC_003197 f	Salmone 3 -- - ----- -
3678	an	CACTGTGG	GCTG	rom 1942	CAA CC TGC GCTG A
		TTTGTA		01 to 19499	GTT GG ACG CGAC
				5 (-)	AT T TGTCAC A
GAM35	Hum	CAGCAGCA	CAACCTGC	spe NC_003198 f	Salmone 3 -- - ----- -
3678	an	CACTGTGG	GCTG	rom 1963	CAA CC TGC GCTG A
		TTTGTA		89 to 19718	GTT GG ACG CGAC
				3 (-)	AT T TGTCAC A
GAM35	Hum	CAGCAGCA	CAACCTGC	spe NC_004631 f	Salmone 3 -- - ----- -
3678	an	CACTGTGG	GCTG	rom 1963	CAA CC TGC GCTG A
		TTTGTA		80 to 19717	GTT GG ACG CGAC
				4 (-)	AT T TGTCAC A
GAM35	Hum	CAGCAGCA	CAACGCCA	flh NC_002929 f	Bordete 3 -- CG -----A CAA
3678	an	CACTGTGG	AGCTGCTG	rom 14417	CCA GCTGCTG GTT A
		TTTGTA		67 to 14429	GGT CGACGAC AT
				21 (+)	T- GTCACA
GAM35	Hum	CAGCAGCA	CAAGCCAA	sse NC_004431 f	Escheri 3 -- A---- -
3678	an	CACTGTGG	TCTGCTG	rom 29224	CAAGCCA T CTGCTG A
		TTTGTA		56 to 29232	GTTTGGT A GACGAC
				41 (-)	AT GTCAC C
GAM35	Hum	CAGCAGCA	CAAGCCCG	aer NC_002947 f	Pseudom 3 -- CGCT-- -
3678	an	CACTGTGG	CTTGCTGT	rom 24069	CAAGCC TGCTG TG A
		TTTGTA		96 to 24085	GTTTGG ACGAC AC
				61 (-)	AT TGTCAC G

GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAAGCCTG CGCTG	lpp I	NC_000962 f rom 22912 67 to 22919 23 (+)	Mycobac teriu m tubercu los is H37Rv	3 --	-----	-	CAAGCC GTTTGG AT	TGC GCTG ACG CGAC TGTCAC A	A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAAGCCTG CGCTG	lpp I	NC_002945 f rom 22751 82 to 22758 38 (+)	Mycobac teriu m bovis subs p bovis AF21 22/97	3 --	-----	-	CAAGCC GTTTGG AT	TGC GCTG ACG CGAC TGTCAC A	A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAAGCTGC TGCTG	ris A	NC_002929 f rom 37652 57 to 37659 91 (-)	Bordete lla p ertussi s	3 --	-----	CAAGC	TGCTGCTG ACGACGAC GIGTCAC	GTTTG AT	A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAAGGACC ATGCGCTG	acc C	NC_002929 f rom 9264 07 to 92777 7 (+)	Bordete lla p ertussi s	3 --	GG	-----	CAA ACCA GCTG GTT TGGT ACG CGAC AT GTCAC A	TGC TGGT AT	A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAAGGCAA AGGICTGC TG	rps T	NC_002947 f rom 7070 68 to 70734 6 (-)	Pseudom onas putida KT244 0	3 --	G A G- -	CAAG	CA AG T CTGCTG GT TC A GACGAC AT G AC C	GTTT G	A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAATAACA ATGCAGCT	fmt B(m r p)	NC_002745 f rom 22181 45 to 22255 90 (-)	Staphyl ococc us aureus su bsp. aureus N315	3 --	T A A----	A	CAA A CA GTT T GT AT - G	TGC GCTG ACG CGAC GTCAC A	A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAATAACA ATGCAGCT	fmt B(m r p)	NC_002758 f rom 22879 35 to 22953 80 (-)	Staphyl ococc us aureus su bsp. aureus Mu50	3 --	T A A----	A	CAA A CA GTT T GT AT - G	TGC GCTG ACG CGAC GTCAC A	A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAATAACA ATGCAGCT	tru nca t ed fmt B	NC_003923 f rom 22380 83 to 22401 43 (-)	Staphyl ococc us aureus su bsp. aureus MW2	3 --	T A A----	A	CAA A CA GTT T GT AT - G	TGC GCTG ACG CGAC GTCAC A	A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAATAGCG CCTGCTGC TG	nup C	NC_004337 f rom 25158 42 to 25170 83 (+)	Shigell a fle xneri 2a str . 301	3 --	T-- GC CC	CAA	A G TGCTGCTG T C ACGACGAC AT TGG GT AC	GTT AT	A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAATAGCG CCTGCTGC TG	nup C	NC_004431 f rom 27953 90 to 27966 31 (+)	Escheri chia coli CFT073	3 --	T-- GC CC	CAA	A G TGCTGCTG T C ACGACGAC AT TGG GT AC	GTT AT	A
GAM35 3678	Hum an	CAGCAGCA CACTGTGG TTTGTA	CAATAGCG CCTGCTGC TG	nup C	NC_004741 f rom 24940 19 to 24952 21 (+)	Shigell a fle xneri 2a str . 2457T	3 --	T-- GC CC	CAA	A G TGCTGCTG T C ACGACGAC AT TGG GT AC	GTT AT	A

GAM35	Hum	CAGCAGCA	CAATATAG	NC_000922 f	Chlamyd				
3678	an	CACTGTGG	AAGCTGCT	def rom 12217	ophil a 3	--	TA G AGC	A	A
		TTTGTA	GCTA	35 to 12222	pneumon		CAA TA A TGCTGCT		
				95 (+)	iae		GTT GT T ACGACGA		
					CWL029		AT TG G CAC		C
GAM35	Hum	CAGCAGCA	CAATATAG	NC_002491 f	Chlamyd				
3678	an	CACTGTGG	AAGCTGCT	def rom 12180	ophil a 3	--	TA G AGC	A	A
		TTTGTA	GCTA	69 to 12186	pneumon		CAA TA A TGCTGCT		
				29 (+)	iae		GTT GT T ACGACGA		
					J138		AT TG G CAC		C
GAM35	Hum	CAGCAGCA	CAATCACC	NC_004431 f	Escheri				
3678	an	CACTGTGG	GGGCCGAT	glc rom 35428	chia 3	--	T - C G CCGA	G	
		TTTGTA	GCGGCTG	C rom 35436	coli		CAA C AC G G TGC	A	
				95 (+)	CFT073		GCTG GTT G TG C C		
							ACG CGAC AT T G T		
							A ---- A		
GAM35	Hum	CAGCAGCA	CAATCAGG	NC_002947 f	Pseudom				
3678	an	CACTGTGG	GATACTGC	pta rom 8916	onas 3	--	T - G A- A CAA	A	
		TTTGTA	TG	25 to 89371	putida		C A GG T CTGCTG GTT		
				2 (-)	KT244 0		G T TC A GACGAC AT T		
							G G AC C		
GAM35	Hum	CAGCAGCA	CAATCCCC	NC_002929 f	Bordete				
3678	an	CACTGTGG	GCTTCCTG	tcf rom 12644	lla p 3	--	T C C - C CAA	A	
		TTTGTA	CTG	A 36 to 12663	ertussi		CC CG T T CTGCTG GTT		
				79 (+)	s		GG GT A A GACGAC AT T		
							T C C C		
GAM35	Hum	CAGCAGCA	CAATCCCG	NC_003143 f	Yersini				
3678	an	CACTGTGG	GCCATTTG	ruv rom 23364	3	--	T CG CCATT -		
		TTTGTA	CTCTG	B 49 to 23374	a pes		CAA CC G TGCT	A	
				53 (+)	tis		CTG GTT GG T		
							ACGA GAC AT T TG		
							CAC-- C		
GAM35	Hum	CAGCAGCA	CAATCCCG	NC_004088 f	Yersini				
3678	an	CACTGTGG	GCCATTTG	ruv rom 24820	3	--	T CG CCATT -		
		TTTGTA	CTCTG	B 31 to 24830	a pes		CAA CC G TGCT	A	
				35 (-)	tis KIM		CTG GTT GG T		
							ACGA GAC AT T TG		
							CAC-- C		
GAM35	Hum	CAGCAGCA	CAATCGCA	NC_004337 f	Shigell				
3678	an	CACTGTGG	GCACTGGT	nup rom 25158	3	--	T G GCAC- G		
		TTTGTA	GCTG	C 42 to 25170	a fle		CAA C CA TG TGCTG	A	
				83 (+)	xneri		GTT G GT AC ACGAC		
					2a str		AT T - GTCAC G		
					. 301				
GAM35	Hum	CAGCAGCA	CAATCTCA	NC_002947 f	Pseudom				
3678	an	CACTGTGG	CTTTCTGC	rps rom 7070	3	--	T T TT C -		
		TTTGTA	GCTG	T 68 to 70734	onas		CAA C CAC T TGC GCTG	A	
				6 (-)	putida		GTT G GTG A ACG CGAC		
					KT244 0		AT T - TC C A		
GAM35	Hum	CAGCAGCA	CACACAGG	NC_002677 f	Mycobac				
3678	an	CACTGTGG	TCATCCTT	prc rom 15765	3	--	C - G TCATCCT -		
		TTTGTA	GCGCTG	A 53 to 15773	teriu m		CA AC A G TGC	A	
				50 (+)	leprae		GCTG GT TG T T		
							ACG CGAC AT T G		
							G CAC---- A		
GAM35	Hum	CAGCAGCA	CACATGTT	NC_004337 f	Shigell				
3678	an	CACTGTGG	GTACATGC	nup rom 25158	3	--	- - - T TACA		
		TTTGTA	TGCTT	C 42 to 25170	a fle		T CA C ATG TG	A	
				83 (+)	xneri		TGCTGCT GT G TGT		
					2a str		AC ACGACGA AT TT G		
					. 301		C ---- C		
GAM35	Hum	CAGCAGCA	CACCAATG	NC_000962 f	Mycobac				
3678	an	CACTGTGG	CTCCTG	lpp rom 22912	3	--	- - A---- C		
		TTTGTA		I 67 to 22919	teriu m		CA CCA TGCT CTG	A	
				23 (+)	tubercu		GT GGT ACGA GAC		
					los is		AT TT GTCAC C		
					H37Rv				
GAM35	Hum	CAGCAGCA	CACCAATG	NC_002945 f	Mycobac				
3678	an	CACTGTGG	CTCCTG	lpp rom 22751	3	--	- - A---- C		
		TTTGTA		I 82 to 22758	teriu m		CA CCA TGCT CTG	A	
					bovis		GT GGT ACGA GAC		

				38	(+)	subs p	AT	TT	GTCAC	C	
						bovis					
						AF21					
						22/97					
GAM35	Hum	CAGCAGCA	CACCACCG	omp	NC_004431 f	Escheri	3	--	--	C CTAAC	
3678	an	CACGTGTG	CTAACTGC	G	rom 16245	chia			-	CA CCAC G	A
		TTTGTA	TGCG		77 to 16255	coli			TGCTGC G	GT GGTG	
					33 (+)	CFT073		C	ACGACG C	AT TT	
									T AC---	A	
						Salmone					
GAM35	Hum	CAGCAGCA	CACCACCT	phn	NC_003198 f	lla e	3	--	--	C--- C	CA
3678	an	CACGTGTG	CCTGCTG	V	rom 4715	nterica			CCAC	T CTGCTG	GT
		TTTGTA			75 to 47236	ente			GGTG	A GACGAC	AT
					6 (-)	rica			TT	TCAC C	
						serovar					
						Typhi					
						Salmone					
GAM35	Hum	CAGCAGCA	CACCACCT	phn	NC_004631 f	lla e	3	--	--	C--- C	CA
3678	an	CACGTGTG	CCTGCTG	V	rom 25087	nterica			CCAC	T CTGCTG	GT
		TTTGTA			35 to 25095	ente			GGTG	A GACGAC	AT
					26 (+)	rica			TT	TCAC C	
						serovar					
						Typhi					
						Ty2					
GAM35	Hum	CAGCAGCA	CACCACGT	pil	NC_002947 f	Pseudom	3	--	--	TA	T
3678	an	CACGTGTG	AGTGCTTC	T	rom 58169	onas			CA	CCACG	GTGCT CTG
		TTTGTA	TG		34 to 58179	putida			GT	GGTGT	CACGA GAC
					44 (-)	KT244 0		AT	TT	CA	C
GAM35	Hum	CAGCAGCA	CACCAGCG	oxy	NC_003197 f	Salmone	3	--	--	GC CC	- CA
3678	an	CACGTGTG	CCTGCGCT	G R	rom 43430	lla t			CCA	G TGC	GCTG GT
		TTTGTA			80 to 43439	yphimur			GGT	C ACG	CGAC AT
					97 (+)	ium L			TT	GT AC	A
						T2					
						Salmone					
GAM35	Hum	CAGCAGCA	CACCAGCG	oxy	NC_003198 f	lla e	3	--	--	GC CC	- CA
3678	an	CACGTGTG	CCTGCGCT	G R	rom 36072	nterica			CCA	G TGC	GCTG GT
		TTTGTA			04 to 36081	ente			GGT	C ACG	CGAC AT
					21 (-)	rica			TT	GT AC	A
						serovar					
						Typhi					
						Salmone					
GAM35	Hum	CAGCAGCA	CACCAGCG	oxy	NC_004631 f	lla e	3	--	--	GC CC	- CA
3678	an	CACGTGTG	CCTGCGCT	G R	rom 35928	nterica			CCA	G TGC	GCTG GT
		TTTGTA			64 to 35937	ente			GGT	C ACG	CGAC AT
					81 (-)	rica			TT	GT AC	A
						serovar					
						Typhi					
						Ty2					
GAM35	Hum	CAGCAGCA	CACCATGC	MGA	NM_002410	Human	3	--	--	-----	CA
3678	an	CACGTGTG	TGCTG	T5					CCA	TGCTGCTG	GT
		TTTGTA							GGT	ACGACGAC	AT
								TT	GTCAC		
						Shigell					
GAM35	Hum	CAGCAGCA	CACCATTG	ipa	NC_004741 f	a file	3	--	--	-----	C
3678	an	CACGTGTG	CTGCCG	H_5	rom 20232	xneri			CA	CCAT	TGCTGC G
		TTTGTA			05 to 20248	2a str			GT	GGTG	ACGACG C
					48 (+)	. 2457T		AT	TT	TCAC	A
GAM35	Hum	CAGCAGCA	CACCATTG	ipa	NC_004337 f	Shigell	3	--	--	-----	C
3678	an	CACGTGTG	CTGCCG	H9.	rom 14220	a file			CA	CCAT	TGCTGC G
		TTTGTA		8	64 to 14237	xneri			GT	GGTG	ACGACG C
					79 (-)	2a str		AT	TT	TCAC	A

				. 301								
GAM35	Hum	CAGCAGCA			NC_004337 f	Shigell	3	--	--	----	C	A
3678	an	CACGTGTG	CACCATTTG	sit	rom 14053	a file				CA CCAT	TGCTGC	G
		TTTGTA	CTGCCG	D	60 to 14062	xneri				GT GGTG	ACGACG	C
					17 (-)	2a str				AT TT	TCAC	A
					. 301							
						Shigell						
GAM35	Hum	CAGCAGCA			NC_004741 f	a file	3	--	--	----	C	A
3678	an	CACGTGTG	CACCATTTG	sit	rom 19046	xneri				CA CCAT	TGCTGC	G
		TTTGTA	CTGCCG	D	66 to 19055	2a str				GT GGTG	ACGACG	C
					23 (+)	. 2457T				AT TT	TCAC	A
GAM35	Hum	CAGCAGCA			NC_002677 f	Mycobac	3	--	--	TTC-	C	CA
3678	an	CACGTGTG	CACCATTT	dna	rom 14230	teriu m				CCAT	TGC GCTG	GT
		TTTGTA	CTGCCGCT	G E	14 to 14265	leprae				GGTG	ACG CGAC	AT
					47 (+)					TT	TCAC	A
GAM35	Hum	CAGCAGCA			NC_002677 f	Mycobac	3	--	--	TTC-	C	CA
3678	an	CACGTGTG	CACCATTT	pcn	rom 32482	teriu m				CCAT	TGC GCTG	GT
		TTTGTA	CTGCCGCT	G A	68 to 32497	leprae				GGTG	ACG CGAC	AT
					28 (-)					TT	TCAC	A
GAM35	Hum	CAGCAGCA			NC_002677 f	Mycobac	3	--	--	TTC-	C	CA
3678	an	CACGTGTG	CACCATTT	pol	rom 16482	teriu m				CCAT	TGC GCTG	GT
		TTTGTA	CTGCCGCT	G A	20 to 16509	leprae				GGTG	ACG CGAC	AT
					55 (-)					TT	TCAC	A
GAM35	Hum	CAGCAGCA			NC_002677 f	Mycobac	3	--	--	TTC-	C	CA
3678	an	CACGTGTG	CACCATTT	tru	rom 23433	teriu m				CCAT	TGC GCTG	GT
		TTTGTA	CTGCCGCT	G A	29 to 23440	leprae				GGTG	ACG CGAC	AT
					78 (-)					TT	TCAC	A
GAM35	Hum	CAGCAGCA			NC_004431 f	Escheri	3	--	C-	GC	T	T
3678	an	CACGTGTG	CACCCAGC	dad	rom 14763	chia				CA CCA	GT TGCTGCT	A
		TTTGTA	GTTTGCTG	X	06 to 14773	coli				GT GGT	CA ACGACGA	
			CTT		76 (+)	CFT073				AT TT	GT C	C
GAM35	Hum	CAGCAGCA			NC_000962 f	Mycobac	3	--	C-	G	TGTC	CA
3678	an	CACGTGTG	CACCCAGG	nic	rom 31666	teriu m				CCA G	GCTGCTG	GT
		TTTGTA	TGTCGCTG	T	81 to 31677	tubercu				GGT T	CGACGAC	AT
			CTG		99 (+)	los is				TT	G CACA	
						H37Rv						
						Mycobac						
GAM35	Hum	CAGCAGCA			NC_002945 f	teriu m	3	--	C-	G	TGTC	CA
3678	an	CACGTGTG	CACCCAGG	nic	rom 31232	bovis				CCA G	GCTGCTG	GT
		TTTGTA	TGTCGCTG	T	00 to 31243	subs p				GGT T	CGACGAC	AT
			CTG		18 (+)	bovis				TT	G CACA	
						AF21						
						22/97						
GAM35	Hum	CAGCAGCA			NC_002929 f	Bordete	3	--	--	T TC--		C
3678	an	CACGTGTG	CACCTTTC	ris	rom 37652	lla p				CA CC T	TGCTGCT	A
		TTTGTA	TGCTGCTC	A	57 to 37659	ertussi				GT GG G	ACGACGA	
					91 (-)	s				AT TT	T TCAC	C
GAM35	Hum	CAGCAGCA			NC_004741 f	Shigell	3	--	C		TGTGTT	-
3678	an	CACGTGTG	CACGCCAT	ipa	rom 20232	a file				CA GCCATA	TG	A
		TTTGTA	ATGTGTTT	H_5	05 to 20248	xneri				TGCTG	GT TGGTGT	
			GTGCTG		48 (+)	2a str				AC ACGAC	AT T	
						. 2457T				CAC---	G	
GAM35	Hum	CAGCAGCA			NC_004337 f	Shigell	3	--	C		TGTGTT	-
3678	an	CACGTGTG	CACGCCAT	ipa	rom 14220	a file				CA GCCATA	TG	A
		TTTGTA	ATGTGTTT	H9.	64 to 14237	xneri				TGCTG	GT TGGTGT	
			GTGCTG	8	79 (-)	2a str				AC ACGAC	AT T	
						. 301				CAC---	G	
GAM35	Hum	CAGCAGCA			NC_004337 f	Shigell	3	--	C		TGTGTT	-
3678	an	CACGTGTG	CACGCCAT	sit	rom 14053	a file				CA GCCATA	TG	A
			ATGTGTTT	D								

	TTTGTA	GTGCTG		60 to 14062	xneri	TGCTG	GT TGGTGT	
				17 (-)	2a str		AC ACGAC AT T	
					. 301		CAC--- G	
					Shigell			
GAM35	Hum	CAGCAGCA	CACGCCAT	sit	NC_004741 f	Shigell	3	-- C TGTT -
3678	an	CACGTGG	ATGTGTT	D	rom 19046	a file	3	CA GCCATA TG A
		TTTGTA	GTGCTG		66 to 19055	xneri		TGCTG GT TGGTGT
					23 (+)	2a str		AC ACGAC AT T
						. 2457T		CAC--- G
						Shigell		
GAM35	Hum	CAGCAGCA	CAGAACAC	rb	NC_004337 f	Shigell	3	-- A T TACC
3678	an	CACGTGG	TGTTACCG	R	rom 39477	a file	3	CAGA CAC GT
		TTTGTA	CTGCTG		08 to 39487	xneri		GCTGCTG GTTT GTG
					00 (+)	2a str		CA CGACGAC AT G
						. 301		T C--A
						Escheri		
GAM35	Hum	CAGCAGCA	CAGAACAC	rb	NC_004431 f	Escheri	3	-- A T TACC
3678	an	CACGTGG	TGTTACCG	R	rom 44392	chia	3	CAGA CAC GT
		TTTGTA	CTGCTG		60 to 44402	coli		GCTGCTG GTTT GTG
					52 (+)	CFT073		CA CGACGAC AT G
								T C--A
						Shigell		
GAM35	Hum	CAGCAGCA	CAGAACAC	rb	NC_004741 f	Shigell	3	-- A T TACC
3678	an	CACGTGG	TGTTACCG	R	rom 38245	a file	3	CAGA CAC GT
		TTTGTA	CTGCTG		94 to 38255	xneri		GCTGCTG GTTT GTG
					77 (-)	2a str		CA CGACGAC AT G
						. 2457T		T C--A
						Shigell		
GAM35	Hum	CAGCAGCA	CAGAAGCA	rb	NC_004337 f	Shigell	3	-- G G AACAT CA
3678	an	CACGTGG	TGAACATT	R	rom 39477	a file	3	AA CATG TGCTGCTG A
		TTTGTA	GCTGCTG		08 to 39487	xneri		GT TT GTGT
					00 (+)	2a str		ACGACGAC AT - G
						. 301		CAC--
						Shigell		
GAM35	Hum	CAGCAGCA	CAGAAGCA	rb	NC_004741 f	Shigell	3	-- G G AACAT CA
3678	an	CACGTGG	TGAACATT	R	rom 38245	a file	3	AA CATG TGCTGCTG A
		TTTGTA	GCTGCTG		94 to 38255	xneri		GT TT GTGT
					77 (-)	2a str		ACGACGAC AT - G
						. 2457T		CAC--
						Pseudom		
GAM35	Hum	CAGCAGCA	CAGACGAT	fep	NC_002516 f	Pseudom	3	-- G C--- C
3678	an	CACGTGG	CTCCTGCT	G C	rom 46535	onas		CAGAC AT T CTGCTG A
		TTTGTA			08 to 46543	aerugin		GTITG TG A GACGAC
					05 (-)	osa P		AT G TCAC C
						A01		
						Pseudom		
GAM35	Hum	CAGCAGCA	CAGACTCA	pil	NC_002947 f	Pseudom	3	-- T GC--- C
3678	an	CACGTGG	GCTGCTGC	T	rom 58169	onas		CAGAC CA TGCTGCT A
		TTTGTA	TC		34 to 58179	putida		GTITG GT ACGACGA
					44 (-)	KT244 0		AT - GTCAC C
						Salmone		
						lla e		
GAM35	Hum	CAGCAGCA	CAGCAGGC	cII	NC_003198 f	Salmone	3	-- - - G C T CAG C
3678	an	CACGTGG	TTTGCTGC		rom 45373	nterica		A G T TGCTGCTG GTT G A
		TTTGTA	TG		12 to 45375	ente		T T A ACGACGAC AT T G
					33 (+)	rica		G C C
						serovar		
						Typhi		
						Salmone		
						lla e		
GAM35	Hum	CAGCAGCA	CAGCAGGC	cII	NC_004631 f	Salmone	3	-- - - G C T CAG C
3678	an	CACGTGG	TTTGCTGC		rom 45201	nterica		A G T TGCTGCTG GTT G A
		TTTGTA	TG		21 to 45203	ente		T T A ACGACGAC AT T G
					42 (+)	rica		G C C
						serovar		
						Typhi		
						Ty2		
GAM35	Hum	CAGCAGCA	CAGCCACA	gad	NC_002947 f	Pseudom	3	-- - ----A CAG A
3678	an	CACGTGG	GCTGCTG		rom 48716	onas		CCAC GCTGCTG GTT

		TTTGTA		25 to 48729 29 (+)	putida KT244 0	GGTG CGACGAC AT T TCACA
Salmone						
GAM35	Hum	CAGCAGCA	CAGCCAGG	phn NC_003198 f	lla e	3 -- - G T T - A
3678	an	CACTGTGG	TTTTGCTC	rom 4715	nterica	CAG CCA G T TGCT CTG
		TTTGTA	TG	75 to 47236	ente	GTT GGT T A ACGA GAC
				6 (-)	rica	AT T G C C C
					serovar	
					Typhi	
Salmone						
GAM35	Hum	CAGCAGCA	CAGCCAGG	phn NC_004631 f	lla e	3 -- - G T T - A
3678	an	CACTGTGG	TTTTGCTC	rom 25087	nterica	CAG CCA G T TGCT CTG
		TTTGTA	TG	35 to 25095	ente	GTT GGT T A ACGA GAC
				26 (+)	rica	AT T G C C C
					serovar	
					Typhi	
					Ty2	
GAM35	Hum	CAGCAGCA	CAGCGAGG	tcf NC_002929 f	Bordete	3 -- - G G CCTCG CAG
3678	an	CACTGTGG	CCTCGTGC	rom 12644	lla p	C A G TGCTGCTG A
		TTTGTA	TGCTG	36 to 12663	ertussi	GTT G T T
				79 (+)	s	ACGACGAC AT T G G
						CAC--
GAM35	Hum	CAGCAGCA	CAGCGCTT	uhp NC_003143 f	Yersini	3 -- CG T A CAG
3678	an	CACTGTGG	GGTGATGC	rom 45227	a pes	C TGGTG TGCTGCTG GTT A
		TTTGTA	TGCTG	90 to 45233	tis	G GTCAC ACGACGAC AT
				80 (-)		TG T -
GAM35	Hum	CAGCAGCA	CAGGCGCA	cys NC_002947 f	Pseudom	3 -- G G CAGGC CA A
3678	an	CACTGTGG	GGGTGTGC	rom 3151	onas	GGTGTGCTGCTG GTTTG GT
		TTTGTA	TGCTG	84 to 31598	putida	TCACACGACGAC AT -
				4 (+)	KT244 0	G
GAM35	Hum	CAGCAGCA	CATACCTC	pbp NC_002947 f	Pseudom	3 -- T T CCGCAC C
3678	an	CACTGTGG	CCGCACTG	rom 43237	onas	CA ACC C TGCTGC A
		TTTGTA	CTGCCG	07 to 43246	putida	G GT TGG G
				33 (+)	KT244 0	ACGACG C AT T T
						TCAC-- A
GAM35	Hum	CAGCAGCA	CATAICTG	ung NC_000907 f	Haemoph	3 -- T ----- CA ATC A
3678	an	CACTGTGG	CTGCTG	rom 186	ilus	TGCTGCTG GT TGG
		TTTGTA		76 to 19335	influen	ACGACGAC AT T
				(+)	zae R d	TGTCAC
GAM35	Hum	CAGCAGCA	CATCCACA	ssb NC_002947 f	Pseudom	3 -- T- ---C CA A
3678	an	CACTGTGG	CGCTGCTG	rom 5710	onas	CCACA GCTGCTG GT
		TTTGTA		27 to 57157	putida	GGTGT CGACGAC AT
				2 (+)	KT244 0	TT CACA
GAM35	Hum	CAGCAGCA	CATCCATA	rbs NC_004431 f	Escheri	3 -- T- TC CCAT
3678	an	CACTGTGG	TCGCCATT	rom 44392	chia	G CA CCATA G A
		TTTGTA	GCTGGTG	60 to 44402	coli	TGCTG TG GT GGTGT
				52 (+)	CFT073	C ACGAC AC AT TT
						CA ---- G
GAM35	Hum	CAGCAGCA	CATCGCGG	glp NC_003143 f	Yersini	3 -- TCG G CGCGC
3678	an	CACTGTGG	GCGCGCTG	rom 42896	a pes	C CA CG G A
		TTTGTA	CTGCTC	50 to 42908	tis	TGCTGCT GT GT T
				97 (-)		ACGACGA AT TTG G
						CAC-- C
GAM35	Hum	CAGCAGCA	CATCGCGG	glp NC_004088 f	Yersini	3 -- TCG G CGCGC
3678	an	CACTGTGG	GCGCGCTG	rom 4546	a pes	C CA CG G A
		TTTGTA	CTGCTC	77 to 45604	tis KIM	TGCTGCT GT GT T
				7 (+)		ACGACGA AT TTG G
						CAC-- C
GAM35	Hum	CAGCAGCA	CATGTCGG	dna NC_002677 f	Mycobac	3 -- T GT GTGGA A

3678	an	CACGTGTGG TTTGTA	TGGTGGAT GCTGCTT	E	rom 14230 14 to 14265 47 (+)	teriu m leprae	T CA GTCG G TGCTGCT GT TGGT C ACGACGA AT T GT AC--- C	
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CATGTCGG TGGTGGAT GCTGCTT	pcn A	NC_002677 f rom 32482 68 to 32497 28 (-)	Mycobac 3 teriu m leprae	-- T GT GTGGA T CA GTCG G TGCTGCT GT TGGT C ACGACGA AT T GT AC--- C	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CATGTCGG TGGTGGGT GCTGCTT	pol A	NC_002677 f rom 16482 20 to 16509 55 (-)	Mycobac 3 teriu m leprae	-- T GT GTGGG T CA GTCG G TGCTGCT GT TGGT C ACGACGA AT T GT AC--- C	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CATGTCGG TGGTGGGT GCTGCTT	tru A	NC_002677 f rom 23433 29 to 23440 78 (-)	Mycobac 3 teriu m leprae	-- T GT GTGGG T CA GTCG G TGCTGCT GT TGGT C ACGACGA AT T GT AC--- C	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CCCCTGCT GCTT	omp G	NC_004431 f rom 16245 77 to 16255 33 (+)	Escheri 3 chia coli CFT073	-- --- ----- T C CCG TGCTGCT G GGT ACGACGA AT TTT GTCAC C	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CCCTCGGT GCTGCTG	flh B	NC_002929 f rom 14417 67 to 14429 21 (+)	Bordete 3 lla p ertussi s	-- --- T -- C CC CGG TGCTGCTG G GG GTC ACGACGAC AT TTT T AC	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CGAACCAC CGATGCTG TG	aer -2	NC_002947 f rom 24069 96 to 24085 61 (-)	Pseudom 3 onas putida KT244 0	-- C A- - CGAACCAC G TGCTG TG GTTTGGTG C ACGAC AC AT T AC G	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CGAACGTG CTGCTG	prc A	NC_002677 f rom 15765 53 to 15773 50 (+)	Mycobac 3 teriu m leprae	-- G----- CGAAC TGCTGCTG GTTTG ACGACGAC AT GTGTCAC	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CGACCACC GCGTGGTG CTG	van B	NC_002516 f rom 55041 20 to 55050 73 (+)	Pseudom 3 onas aerugin osa P A01	-- - C C G CGA CCAC G GTG TGCTG GTT GGTG C CAC ACGAC AT T T A G	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CGACTGCT GCTG	pch A	NC_002516 f rom 47451 20 to 47465 50 (+)	Pseudom 3 onas aerugin osa P A01	-- - ----- CGA C TGCTGCTG GTT G ACGACGAC AT T GTGTCAC	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CGAGCGAT GCTGCTT	dsd A	NC_003197 f rom 40044 53 to 40057 75 (+)	Salmone 3 lla t yphimur ium L T2	-- G ----- T CGAGC A TGCTGCT GTTG T ACGACGA AT G GTCAC C	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CGAGGGAT GTAGTGCT GCTC	nic T	NC_000962 f rom 31666 81 to 31677 99 (+)	Mycobac 3 teriu m tubercu los is H37Rv	-- GG TA C CGAG ATG GTGCTGCT GTTT TGT CACGACGA AT GG CA C	A
GAM35 3678	Hum an	CAGCAGCA CACGTGTGG TTTGTA	CGAGGGAT GTAGTGCT GCTC	nic T	NC_002945 f rom 31232 00 to 31243 18 (+)	Mycobac teriu m 3 bovis subs p bovis AF21	-- GG TA C CGAG ATG GTGCTGCT GTTT TGT CACGACGA AT GG CA C	A

-25-

3678	an	CACGTGG TTTGTA	TATCTGCT G	rom 45201 21 to 45203 42 (+)	lla e nterica ente rica serovar Typhi Ty2	AACC T CTGCTG G TTGG A GACGAC AT T TGTCAC C
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CTACCTGC CTGCTGCT G A	uhp NC_003143 f rom 45227 90 to 45233 80 (-)	Yersini 3 a pes tis	-- T - TGCC-- C A CC A TGCTGCTG G T GG ACGACGAC AT T T TGTCAC
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CTAGCCCT GCTGCTC	aco NC_002516 f rom 46395 01 to 46413 78 (-)	Pseudom 3 onas aerugin osa P A01	-- T C----- C A C AGCC TGCTGCT C G TTGG ACGACGA AT T TGTCAC C
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CTCCTGCT GCTG	ssb NC_002947 f rom 5710 27 to 57157 2 (+)	Pseudom 3 onas putida KT244 0	-- T-- ----- C CC A TGCTGCTG G GG ACGACGAC AT TTT TGTCAC
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CTCGGGTT TCTTGCTG CTG	fha NC_002929 f rom 30858 65 to 30984 55 (+)	Bordete 3 lla p ertussi s	-- T-- G GTT CT C C A G T TGCTGCTG G G T A ACGACGAC AT TTT G GTC C-
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CTGACCTT GCTGCTC	pta NC_002947 f rom 8916 25 to 89371 2 (-)	Pseudom 3 onas putida KT244 0	-- T T----- C A C GACC TGCTGCT C G TTGG ACGACGA AT T TGTCAC C
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CTGCCTGC TGCTC	orn NC_003143 f rom 3783 31 to 37887 6 (+)	Yersini 3 a pes tis	-- T - ----- C A C G CC TGCTGCT C G T GG ACGACGA AT T T TGTCAC C
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CTGCGTGC TGCTT	glp NC_004310 f rom 2107 63 to 21227 4 (+)	Brucell 3 a sui s 1330	-- T - G----- T A C G C TGCTGCT C G T G ACGACGA AT T T GTGTCAC C
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CTTACTTG CTGCIC	sse NC_004431 f rom 29224 56 to 29232 41 (-)	Escheri 3 chia coli CFT073	-- TT ----- C A C ACT TGCTGCT C G TGG ACGACGA AT TT TGTCAC C
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CTTCTGCTG TGCTT	def NC_000922 f rom 12217 35 to 12222 95 (+)	Chlamyd ophil a 3 pneumon iae CWL029	-- TT- ----- T A C CC TGCTGCT C G GG ACGACGA AT TTT TGTCAC C
GAM35 3678	Hum an	CAGCAGCA CACGTGG TTTGTA	CTTCTGCTG TGCTT	def NC_002491 f rom 12180 69 to 12186 29 (+)	Chlamyd 3 ophil a pneumon iae J138	-- TT- ----- T A C CC TGCTGCT C G GG ACGACGA AT TTT TGTCAC C

Replace paragraph 0159 with the following paragraph.

It is appreciated that the specific functions and accordingly the utilities of each of a plurality of GAM oligonucleotides that are described by Fig. 1 are correlated with and may be deduced from the

identity of the GAM TARGET GENES inhibited thereby, and whose functions are set forth in Table 8, hereby incorporated herein. Table 8, lines 685695-687709 shows data relating to the function and utilities of GAM RNA as set forth in SEQ ID NO: 348.

After paragraph 0159, add the following Table 8.

Table 8

GAM NAME	GAM RNA SEQUENCE	GAM ORGANISM	TARGET GET	TARGET ORGANISM	GAM FUNCTION	GAM POS
=====	=====	=====	=====	=====	=====	=====
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	accC	Bordetella pertussis	GAM353678 is a human miRNA-like oligonucleotide, which targets biotin carboxylase (accC, NC_002929 from 926407 to 927777 (+)), a bacterial target gene encoded by the Bordetella pertussis genome, as part of an anti-bacterial host defense mechanism. accC BINDING SITE 1 and accC BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the accC gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of accC BINDING SITE 1 and accC BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. A function of GAM353678 is to inhibit accC, a bacterial target gene which is associated with Bordetella pertussis infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Bordetella pertussis infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	aceK	Pseudomonas putida KT2440	GAM353678 is a human miRNA-like oligonucleotide, which targets isocitrate dehydrogenase kinase/phosphatase (aceK, NC_002947 from 5184742 to 5186457 (-)), a bacterial target gene encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. aceK BINDING SITE 1 and aceK BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the aceK gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of aceK BINDING SITE 1 and aceK BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit aceK, a bacterial target gene which is associated with Pseudomonas putida KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of	A

					GAM353678 include the diagnosis, prevention and treatment of <i>Pseudomonas putida</i> K T2440 infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	acoR	<i>Pseudomonas aeruginosa</i> PA01	GAM353678 is a human miRNA-like oligonucleotide, which targets transcriptional regulator AcoR (acoR, NC_002516 from 463950 1 to 4641378 (-)), a bacterial target gene encoded by the <i>Pseudomonas aeruginosa</i> PA01 genome, as part of an anti-bacterial host defense mechanism. acoR BINDING SITE 1 and acoR BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the acoR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of acoR BINDING SITE 1 and acoR BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit acoR, a bacterial target gene which is associated with <i>Pseudomonas aeruginosa</i> PA01 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Pseudomonas aeruginosa</i> PA01 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	aer-2	<i>Pseudomonas putida</i> KT2440	GAM353678 is a human miRNA-like oligonucleotide, which targets aerotaxis receptor Aer-2 (aer-2, NC_002947 from 2406996 to 2408561 (-)), a bacterial target gene encoded by the <i>Pseudomonas putida</i> KT2440 genome, as part of an anti-bacterial host defense mechanism. aer-2 BINDING SITE 1 and aer-2 BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the aer-2 gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of aer-2 BINDING SITE 1 and aer-2 BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit aer-2, a bacterial target gene which is associated with <i>Pseudomonas putida</i> KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Pseudomonas putida</i> KT2440 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	amtB	<i>Shigella flexneri</i> 2a str. 301	GAM353678 is a human miRNA-like oligonucleotide, which targets probable ammonium transporter (amtB, NC_004337 from 408059 to 409345 (+)), a bacterial target gene encoded by the <i>Shigella flexneri</i> 2a str. 301 genome, as part of an anti-bacterial host defense mechanism. amtB BINDING SITE 1 and amtB BINDING SITE	A

2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the amtB gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of amtB BINDING SITE 1 and amtB BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit amtB, a GAM353678 bacterial target gene which is associated with *Shigella flexneri* 2a str. 301 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Shigella flexneri* 2a str. 301 infection and associated clinical conditions

GAM35	CAGCAGCA	Human	amtB	Escherichia coli C	GAM353678 is a human miRNA-like	A
3678	CACTGTGG			FT073	oligonucleotide, which targets Probable	
	TTTGTA				ammonium transporter (amtB, NC_004431 from	

547616 to 548902 (+)), a bacterial target gene encoded by the *Escherichia coli* CFT073 genome, as part of an anti-bacterial host defense mechanism. amtB BINDING SITE 1 and amtB BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the amtB gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of amtB BINDING SITE 1 and amtB BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit amtB, a GAM353678 bacterial target gene which is associated with *Escherichia coli* CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Escherichia coli* CFT073 infection and associated clinical conditions

GAM35	CAGCAGCA	Human	amtB	Shigella	GAM353678 is a human miRNA-like	A
3678	CACTGTGG			flexneri	oligonucleotide, which targets probable	
	TTTGTA			2a str. 2	ammonium transporter (amtB, NC_004741 from	
				457T	407860 to 409146 (+)), a bacterial	

target gene encoded by the *Shigella flexneri* 2a str. 2457T genome, as part of an anti-bacterial host defense mechanism. amtB BINDING SITE 1 and amtB BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the amtB gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of amtB BINDING SITE 1 and amtB BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit amtB, a GAM353678

				bacterial target gene which is associated with <i>Shigella flexneri</i> 2a str. 2457T infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Shigella flexneri</i> 2a str. 2457T infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	aroA	<i>Neisseria meningitidis</i> Z2491	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets 5-enolpyruvylshikimate-3-phosphate synthase (aroA, NC_003116 from 1557502 to 1558803 (+)), a bacterial target gene encoded by the <i>Neisseria meningitidis</i> Z2491 genome, as part of an anti-bacterial host defense mechanism. aroA BINDING SITE 1 and aroA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the aroA gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of aroA BINDING SITE 1 and aroA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit aroA, a GAM353678 bacterial target gene which is associated with <i>Neisseria meningitidis</i> Z2491 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Neisseria meningitidis</i> Z2491 infection and associated clinical conditions</p>
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	aroD	<i>Leptospira interrogans</i> serovar lai str. 56601	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets 3-dehydroquinate dehydratase (aroD, NC_004342 from 48128 to 48832 (-)), a bacterial target gene encoded by the <i>Leptospira interrogans</i> serovar lai str. 56601 genome, as part of an anti-bacterial host defense mechanism. aroD BINDING SITE 1 and aroD BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the aroD gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of aroD BINDING SITE 1 and aroD BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit aroD, a GAM353678 bacterial target gene which is associated with <i>Leptospira interrogans</i> serovar lai str. 56601 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Leptospira interrogans</i> serovar lai str. 56601 infection and associated clinical conditions</p>
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	aroH	<i>Shigella flexneri</i> 2a str. 3	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets 3-deoxy-D-arabinoheptulosonate-7-phosphate synthase</p>

01					(DAHP synthetase, tryptophan repressible) (aroH, NC_004337 from 15575 27 to 1558573 (-)), a bacterial target gene encoded by the <i>Shigella flexneri</i> 2a str. 301 genome, as part of an anti-bacterial host defense mechanism. aroH BINDING SITE 1 and aroH BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the aroH gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of aroH BINDING SITE 1 and aroH BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit aroH, a GAM353678 bacterial target gene which is associated with <i>Shigella flexneri</i> 2a str. 301 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Shigella flexneri</i> 2a str. 301 infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	avtA	<i>Shigella flexneri</i> 2a str. 301	GAM353678 is a human miRNA-like oligonucleotide, which targets alanine-alpha-ketoisovalerate (or valine-pyruvate) transaminase, transaminase C (avtA, NC_004337 from 3721175 to 3722533 (+)), a bacterial target gene encoded by the <i>Shigella flexneri</i> 2a str. 301 genome, as part of an anti-bacterial host defense mechanism. avtA BINDING SITE is a bacterial target binding site that is a found in the the 3' untranslated region of mRNA encoded by the avtA gene, corresponding to a target binding site such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of avtA BINDING SITE, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit avtA, a GAM353678 bacterial target gene which is associated with <i>Shigella flexneri</i> 2a str. 301 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Shigella flexneri</i> 2a str. 301 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	avtA	<i>Shigella flexneri</i> 2a str. 2457T	GAM353678 is a human miRNA-like oligonucleotide, which targets alanine-alpha-ketoisovalerate/valine-pyruvate transaminase C (avtA, NC_004741 from 4052685 to 4053938 (-)), a bacterial target gene encoded by the <i>Shigella flexneri</i> 2a str. 2457T genome, as part of an anti-bacterial host defense mechanism. avtA BINDING SITE is a bacterial target binding site that is a found in the the 3' untranslated region of mRNA encoded by the avtA gene, corresponding to a target binding site such as BINDING SITE I, BINDING SITE II or BINDING SITE III of	A

Fig. 1. The nucleotide sequences of avtA BINDING SITE, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit avtA, a GAM353678 bacterial target gene which is associated with *Shigella flexneri* 2a str. 2457T infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Shigella flexneri* 2a str. 2457T infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	cII	Salmonella enterica serovar Typhi Ty2	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets transcriptional regulatory protein (cII, NC_004631 from 452 0121 to 4520342 (+)), a bacterial target gene encoded by the <i>Salmonella enterica</i> serovar Typhi Ty2 genome, as part of an anti-bacterial host defense mechanism. cII BINDING SITE 1 and cII BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the cII gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of cII BINDING SITE 1 and cII BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit cII, a GAM353678 bacterial target gene which is associated with <i>Salmonella enterica</i> serovar Typhi Ty2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella enterica</i> serovar Typhi Ty2 infection and associated clinical conditions</p>	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	cII	Salmonella enterica serovar Typhi	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets transcriptional regulatory protein (cII, NC_003198 from 453 7312 to 4537533 (+)), a bacterial target gene encoded by the <i>Salmonella enterica</i> serovar Typhi genome, as part of an anti-bacterial host defense mechanism. cII BINDING SITE 1 and cII BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the cII gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of cII BINDING SITE 1 and cII BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit cII, a GAM353678 bacterial target gene which is associated with <i>Salmonella enterica</i> serovar Typhi infection, as part of an anti-bacterial host defense mechanism. Accordingly,</p>	A

					the utilities of GAM 353678 include the diagnosis, prevention and treatment of Salm onella enterica enterica serovar Typhi infection and associate d clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	cysM	Salmonella typhimurium LT2	GAM353678 is a human miRNA-like oligonucleotide, which targets cysteine synthase B (cysM, NC_003197 from 2551651 to 255256 2 (-)), a bacterial target gene encoded by the Salmonella typhimurium LT2 genome, as part of an anti-bacterial host defense mechanism. cysM BINDING SITE 1 and cysM BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the cysM gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of cysM BINDING SITE 1 and cysM BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit cysM, a GAM353678 bacterial target gene which is associated with Salmonella typhimurium LT2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Salmonella typhimurium LT2 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	cysQ	Pseudomonas putida KT2440	GAM353678 is a human miRNA-like oligonucleotide, which targets 3'(2'),5'-bisphosphate nucleotidase (cysQ, NC_002947 from 315184 to 315984 (+)), a bacterial target gene encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. cysQ BINDING SITE 1 and cysQ BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the cysQ gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of cysQ BINDING SITE 1 and cysQ BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit cysQ, a GAM353678 bacterial target gene which is associated with Pseudomonas putida KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas putida K T2440 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	dadX	Escherichia coli CFT073	GAM353678 is a human miRNA-like oligonucleotide, which targets Alanine racemase, catabolic (dadX, NC_004431 from 1476306 to 1477376 (+)), a bacterial target gene encoded by the Escherichia coli CFT073 genome, as part of an anti-bacterial host defense mechanism. dadX BINDING SITE 1 and dadX BINDING SITE 2 are bacterial target binding sites that are	A

found in the untranslated regions of mRNA encoded by the dadX gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of dadX BINDING SITE 1 and dadX BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit dadX, a GAM353678 bacterial target gene which is associated with Escherichia coli CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT073 infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	def	Chlamydomonas reinhardtii L029	GAM353678 is a human miRNA-like oligonucleotide, which targets Polypeptide Deformylase (def, NC_000922 from 1221735 to 1222295 (+)), a bacterial target gene encoded by the Chlamydomonas reinhardtii CWL029 genome, as part of an anti-bacterial host defense mechanism. def BINDING SITE 1 and def BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the def gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of def BINDING SITE 1 and def BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit def, a GAM353678 bacterial target gene which is associated with Chlamydomonas reinhardtii CWL029 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Chlamydomonas reinhardtii CWL029 infection and associated clinical conditions.	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	def	Chlamydomonas reinhardtii J138	GAM353678 is a human miRNA-like oligonucleotide, which targets polypeptide deformylase (def, NC_002491 from 1218069 to 1218629 (+)), a bacterial target gene encoded by the Chlamydomonas reinhardtii J138 genome, as part of an anti-bacterial host defense mechanism. def BINDING SITE 1 and def BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the def gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of def BINDING SITE 1 and def BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit def, a GAM353678 bacterial target gene which is associated	A

					with <i>Chlamydomonas reinhardtii</i> J138 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Chlamydomonas reinhardtii</i> J138 infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	dnaE	<i>Mycobacterium leprae</i>	GAM353678 is a human miRNA-like oligonucleotide, which targets DNA polymerase III, [alpha] subunit (dnaE, NC_002677 from 1423014 to 1426547 (+)), a bacterial target gene encoded by the <i>Mycobacterium leprae</i> genome, as part of an anti-bacterial host defense mechanism. dnaE BINDING SITE 1 through dnaE BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the dnaE gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of dnaE BINDING SITE 1 through dnaE BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit dnaE, a GAM353678 bacterial target gene which is associated with <i>Mycobacterium leprae</i> infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Mycobacterium leprae</i> infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	dsdA	<i>Salmonella typhimurium</i> LT2	GAM353678 is a human miRNA-like oligonucleotide, which targets D-serine deaminase (dsdA, NC_003197 from 4004453 to 4005775 (+)), a bacterial target gene encoded by the <i>Salmonella typhimurium</i> LT2 genome, as part of an anti-bacterial host defense mechanism. dsdA BINDING SITE 1 and dsdA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the dsdA gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of dsdA BINDING SITE 1 and dsdA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit dsdA, a GAM353678 bacterial target gene which is associated with <i>Salmonella typhimurium</i> LT2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella typhimurium</i> LT2 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	fepC	<i>Pseudomonas aeruginosa</i> PA01	GAM353678 is a human miRNA-like oligonucleotide, which targets ferric enterobactin transport protein FepC (fepC, NC_002516 from 4653508 to 4654305 (-)), a bacterial target gene encoded by the <i>Pseudomonas aeruginosa</i> PA01 genome, as	A

part of an anti-bacterial host defense mechanism. fepC BINDING SITE 1 and fepC BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the fepC gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of fepC BINDING SITE 1 and fepC BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit fepC, a GAM353678 bacterial target gene which is associated with *Pseudomonas aeruginosa* PA01 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Pseudomonas aeruginosa* PA01 infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	fhaL	Bordetella pertussis	GAM353678 is a human miRNA-like oligonucleotide, which targets adhesin (fhaL, NC_002929 from 3085865 to 3098455 (+)), a bacterial target gene encoded by the <i>Bordetella pertussis</i> genome, as part of an anti-bacterial host defense mechanism. fhaL BINDING SITE 1 and fhaL BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the fhaL gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of fhaL BINDING SITE 1 and fhaL BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit fhaL, a GAM353678 bacterial target gene which is associated with <i>Bordetella pertussis</i> infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Bordetella pertussis</i> infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	flhB	Bordetella pertussis	GAM353678 is a human miRNA-like oligonucleotide, which targets flagellar biosynthetic protein FlhB (flhB, NC_002929 from 1441767 to 1442921 (+)), a bacterial target gene encoded by the <i>Bordetella pertussis</i> genome, as part of an anti-bacterial host defense mechanism. flhB BINDING SITE 1 through flhB BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the flhB gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of flhB BINDING SITE 1 through flhB BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby	A

				incorporated herein. Another function of GAM353678 is to inhibit flhB, a GAM353678 bacterial target gene which is associated with <i>Bordetella pertussis</i> infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Bordetella pertussis</i> infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	fmtB(<i>Staphylococcus aureus</i> subsp. <i>aureus</i> N315	GAM353678 is a human miRNA-like oligonucleotide, which targets FmtB protein (fmtB(mrp), NC_002745 from 2218145 to 2225590 (-)), a bacterial target gene encoded by the <i>Staphylococcus aureus</i> subsp. <i>aureus</i> N315 genome, as part of an anti-bacterial host defense mechanism. fmtB(mrp) BINDING SITE 1 and fmtB(mrp) BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the fmtB(mrp) gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of fmtB(mrp) BINDING SITE 1 and fmtB(mrp) BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit fmtB(mrp), a GAM353678 bacterial target gene which is associated with <i>Staphylococcus aureus</i> subsp. <i>aureus</i> N315 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Staphylococcus aureus</i> subsp. <i>aureus</i> N315 infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	fmtB(<i>Staphylococcus aureus</i> subsp. <i>aureus</i> Mu50	GAM353678 is a human miRNA-like oligonucleotide, which targets FmtB protein (fmtB(mrp), NC_002758 from 2287935 to 2295380 (-)), a bacterial target gene encoded by the <i>Staphylococcus aureus</i> subsp. <i>aureus</i> Mu50 genome, as part of an anti-bacterial host defense mechanism. fmtB(mrp) BINDING SITE 1 and fmtB(mrp) BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the fmtB(mrp) gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of fmtB(mrp) BINDING SITE 1 and fmtB(mrp) BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit fmtB(mrp), a GAM353678 bacterial target gene which is associated with <i>Staphylococcus aureus</i> subsp. <i>aureus</i> Mu50 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Staphylococcus aureus</i> subsp.	A

					aureus Mu50 infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	ftsY	Chlamydo- phila pneu- moniae J138	GAM353678 is a human miRNA-like oligonucleotide, which targets cell division protein ftsY (ftsY, NC_002491 from 1113127 to 1113999 (-)), a bacterial target gene encoded by the Chlamydo-phila pneumoniae J138 genome, as part of an anti-bacterial host defense mechanism. ftsY BINDING SITE 1 and ftsY BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the ftsY gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of ftsY BINDING SITE 1 and ftsY BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ftsY, a bacterial target gene which is associated with Chlamydo-phila pneumoniae J138 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Chlamydo-phila pneumoniae J138 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	ftsY	Chlamydo- phila pneu- moniae CWL029	GAM353678 is a human miRNA-like oligonucleotide, which targets Cell Division Protein FtsY (ftsY, NC_000922 from 1115427 to 1116299 (-)), a bacterial target gene encoded by the Chlamydo-phila pneumoniae CWL029 genome, as part of an anti-bacterial host defense mechanism. ftsY BINDING SITE 1 and ftsY BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the ftsY gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of ftsY BINDING SITE 1 and ftsY BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ftsY, a bacterial target gene which is associated with Chlamydo-phila pneumoniae CWL029 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Chlamydo-phila pneumoniae CWL029 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	gad	Pseudomonas putida KT2440	GAM353678 is a human miRNA-like oligonucleotide, which targets guanine aminohydrolase (gad, NC_002947 from 4871625 to 4872929 (+)), a bacterial target gene encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. gad BINDING SITE 1 and gad BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA	A

encoded by the gad gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE I II of Fig. 1. The nucleotide sequences of gad BINDING SITE 1 and gad BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit gad, a GAM353678 bacterial target gene which is associated with *Pseudomonas putida* KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Pseudomonas putida* KT 2440 infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTGTA	Human	glcC	Escherichia coli CFT073	GAM353678 is a human miRNA-like oligonucleotide, which targets Glc operon transcriptional activator (glcC, NC_004431 from 3542871 to 3543695 (+)), a bacterial target gene encoded by the <i>Escherichia coli</i> CFT073 genome, as part of an anti-bacterial host defense mechanism. glcC BINDING SITE 1 and glcC BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the glcC gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of glcC BINDING SITE 1 and glcC BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glcC, a GAM353678 bacterial target gene which is associated with <i>Escherichia coli</i> CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Escherichia coli</i> CFT073 infection and associated clinical conditions	A
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GAM353678	CAGCAGCA CACTGTGG TTGTA	Human	glgP	Salmonella enterica enterica serovar Typhi Ty2	GAM353678 is a human miRNA-like oligonucleotide, which targets glycogen phosphorylase (glgP, NC_004631 from 4129215 to 4131662 (+)), a bacterial target gene encoded by the <i>Salmonella enterica enterica</i> serovar Typhi Ty2 genome, as part of an anti-bacterial host defense mechanism. glgP BINDING SITE 1 and glgP BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the glgP gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of glgP BINDING SITE 1 and glgP BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glgP, a GAM353678 bacterial target gene which is associated with <i>Salmonella enterica enterica</i> serovar	A
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encoded by the glgP gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of glgP BINDING SITE 1 and glgP BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glgP, a GAM353678 bacterial target gene which is associated with *Salmonella enterica enterica* serovar Typhi Ty2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Salmonella enterica enterica* serovar Typhi Ty2 infection and associated clinical conditions

					<p>Typhi Ty2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella enterica enterica</i> serovar Typhi Ty2 infection and associated clinical conditions</p>		
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	glgP	Salmonella enterica enterica serovar Typhi	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets glycogen phosphorylase (glgP, NC_003198 from 4144568 to 414 7015 (+)), a bacterial target gene encoded by the <i>Salmonella enterica enterica</i> serovar Typhi genome, as part of an anti-bacterial host defense mechanism. glgP BINDING SITE 1 and glgP BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the glgP gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of glgP BINDING SITE 1 and glgP BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glgP, a GAM353678 bacterial target gene which is associated with <i>Salmonella enterica enterica</i> serovar Typhi infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella enterica enterica</i> serovar Typhi infection and associated clinical conditions</p>	A	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	glpC	Yersinia pestis KIM	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets sn-glycerol-3-phosphate dehydrogenase (anaerobic), K-small subunit (glpC, NC_004088 from 454677 to 456047 (+)), a bacterial target gene encoded by the <i>Yersinia pestis</i> KIM genome, as part of an anti-bacterial host defense mechanism. glpC BINDING SITE 1 and glpC BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the glpC gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of glpC BINDING SITE 1 and glpC BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glpC, a GAM353678 bacterial target gene which is associated with <i>Yersinia pestis</i> KIM infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Yersinia pestis</i> KIM infection and associated clinical conditions</p>	A	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	glpC	Yersinia pestis	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets anaerobic glycerol-3-phosphate dehydrogenase subunit</p>	A	

C (glpC, NC_003143 from 4289650 to 4290897 (-)), a bacterial target gene encoded by the *Yersinia pestis* genome, as part of an anti-bacterial host defense mechanism. glpC BINDING SITE 1 and glpC BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the glpC gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of glpC BINDING SITE 1 and glpC BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glpC, a GAM353678 bacterial target gene which is associated with *Yersinia pestis* infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Yersinia pestis* infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	glpD	Brucella suis 1330	GAM353678 is a human miRNA-like oligonucleotide, which targets glycerol-3-phosphate dehydrogenase (glpD, NC_004310 from 210763 to 212274 (+)), a bacterial target gene encoded by the <i>Brucella suis</i> 1330 genome, as part of an anti-bacterial host defense mechanism. glpD BINDING SITE 1 and glpD BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the glpD gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of glpD BINDING SITE 1 and glpD BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glpD, a GAM353678 bacterial target gene which is associated with <i>Brucella suis</i> 1330 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Brucella suis</i> 1330 infection and associated clinical conditions	A
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GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	ipaH_5	Shigella flexneri 2a str. 2457T	GAM353678 is a human miRNA-like oligonucleotide, which targets invasion plasmid antigen (ipaH_5, NC_004741 from 2023205 to 2024848 (+)), a bacterial target gene encoded by the <i>Shigella flexneri</i> 2a str. 2457T genome, as part of an anti-bacterial host defense mechanism. ipaH_5 BINDING SITE 1 and ipaH_5 BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the ipaH_5 gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of ipaH_5 BINDING SITE 1 and ipaH_5 BINDING SITE 2, and the	A
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					complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ipaH_5, a GAM353678 bacterial target gene which is associated with <i>Shigella flexneri</i> 2a str. 2457T infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Shigella flexneri</i> 2a str. 2457T infection and associated clinical conditions.		
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	ipaH9.8	<i>Shigella flexneri</i> 2a str. 301	GAM353678 is a human miRNA-like oligonucleotide, which targets invasion plasmid antigen (ipaH9.8, NC_004337 from 1422064 to 1423779 (-)), a bacterial target gene encoded by the <i>Shigella flexneri</i> 2a str. 301 genome, as part of an anti-bacterial host defense mechanism. ipaH9.8 BINDING SITE 1 and ipaH9.8 BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the ipaH9.8 gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of ipaH9.8 BINDING SITE 1 and ipaH9.8 BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ipaH9.8, a GAM353678 bacterial target gene which is associated with <i>Shigella flexneri</i> 2a str. 301 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Shigella flexneri</i> 2a str. 301 infection and associated clinical conditions.	A	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	livH	<i>Bordetella pertussis</i>	GAM353678 is a human miRNA-like oligonucleotide, which targets high-affinity branched-chain amino acid transport system permease protein (livH, NC_002929 from 1144729 to 1145607 (+)), a bacterial target gene encoded by the <i>Bordetella pertussis</i> genome, as part of an anti-bacterial host defense mechanism. livH BINDING SITE is a bacterial target binding site that is found in the 3' untranslated region of mRNA encoded by the livH gene, corresponding to a target binding site such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of livH BINDING SITE, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit livH, a GAM353678 bacterial target gene which is associated with <i>Bordetella pertussis</i> infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and	A	

					treatment of <i>Bordetella pertussis</i> infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	lppI	Mycobacterium tuberculosis H37Rv	GAM353678 is a human miRNA-like oligonucleotide, which targets lppI (lppI, NC_000962 from 2291267 to 2291923 (+)), a bacterial target gene encoded by the <i>Mycobacterium tuberculosis</i> H37Rv genome, as part of an anti-bacterial host defense mechanism. lppI BINDING SITE 1 and lppI BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the lppI gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of lppI BINDING SITE 1 and lppI BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit lppI, a GAM353678 bacterial target gene which is associated with <i>Mycobacterium tuberculosis</i> H37Rv infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Mycobacterium tuberculosis</i> H37Rv infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	lppI	Mycobacterium bovis subsp. bovis AF2122/97	GAM353678 is a human miRNA-like oligonucleotide, which targets Probable lipoprotein lppI (lppI, NC_002945 from 2275182 to 2275838 (+)), a bacterial target gene encoded by the <i>Mycobacterium bovis</i> subsp. bovis AF2122/97 genome, as part of an anti-bacterial host defense mechanism. lppI BINDING SITE 1 and lppI BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the lppI gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of lppI BINDING SITE 1 and lppI BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit lppI, a GAM353678 bacterial target gene which is associated with <i>Mycobacterium bovis</i> subsp. bovis AF2122/97 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM 353678 include the diagnosis, prevention and treatment of <i>Mycobacterium bovis</i> subsp. bovis AF2122/97 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	MGAT5	Human	GAM353678 is a human miRNA-like oligonucleotide, which targets a human mannosyl (alpha-1,6-)-glycoprotein beta-1,6-N-acetylglucosaminyltransferase (MGAT5, Accession number: NM_002410) as part of a host response mechanism associated with a <i>Salmonella typhimurium</i> LT2 infection. MGAT5 BINDING SITE is a human target binding site that is a found	A

in the the 3' untranslated region of mRNA encoded by the M GAT5 gene, corresponding to a target binding site such as BIND ING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. Add itionally, using the binding site prediction system of the pre sent invention GAM353678-A binds to sequences on orthologous UTR of rat(NM_023095). The nucleotide sequences of MGAT5 BINDIN G SITE, and the complementary secondary structure to the nucle otide sequence of GAM353678 RNA are set forth in Tables 6-7, h ereby incorporated herein. Another function of GAM353678 is to inhibit MGAT5, a GAM353678 human target gene which encodes an enzyme that catalyzes beta 1-6 branching on N-linked carbohydrates. MGAT5 is associated with Salmonella typhimurium LT2 infection, and therefore GAM35 3678 is associated with the abovementioned infection, as part of a host response mechanism. Accordingly, the utilities of GA M353678 include the diagnosis, prevention and treatment of Sal monella typhimurium LT2 infection and associated clinical cond itions. The function of MGAT5 and its association with various diseases and clinical conditions has been established by previous stu dies, as described hereinabove with reference to GAM3451.

GAM35 3678	CAGCAGCA CACTGTGG TTTGTA	Human	miaA	Chlamydia trachoma tis	GAM353678 is a human miRNA-like oligonucleotide, which targets tRNA isopentenylpyrophosphate transferase (miaA, NC_000117 from 899276 to 900295 (+)), a bacterial target gene encode d by the Chlamydia trachomatis genome, as part of an anti-bact erial host defense mechanism. miaA BINDING SITE 1 and miaA BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the miaA gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of miaA BINDING SITE 1 and miaA BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit miaA, a GAM353678 bacterial target gene which is associated with Chlamydia trach omatis infection, as part of an anti-bacterial host defense me chanism. Accordingly, the utilities of GAM353678 include the d iagnosis, prevention and treatment of Chlamydia trachomatis in fection and associated clinical conditions	A
GAM35 3678	CAGCAGCA CACTGTGG TTTGTA	Human	minE	Pseudomon as putida KT2440	GAM353678 is a human miRNA-like oligonucleotide, which targets cell division topological specificity factor MinE (minE, NC_0 02947 from 1932680 to 1932934 (-)), a bacterial target gen e encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. minE BINDING SITE is a bacterial target binding site that is a found in the the 3' untranslated region	A

of mRNA encoded by the minE gene, corresponding to a target binding site such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of minE BINDING SITE, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit minE, a GAM353678 bacterial target gene which is associated with *Pseudomonas putida* KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Pseudomonas putida* KT2440 infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	nicT	Mycobacterium tuberculosis H37Rv	GAM353678 is a human miRNA-like oligonucleotide, which targets nicT (nicT, NC_000962 from 3166681 to 3167799 (+)), a bacterial target gene encoded by the <i>Mycobacterium tuberculosis</i> H37Rv genome, as part of an anti-bacterial host defense mechanism. nicT BINDING SITE 1 and nicT BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the nicT gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of nicT BINDING SITE 1 and nicT BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit nicT, a GAM353678 bacterial target gene which is associated with <i>Mycobacterium tuberculosis</i> H37Rv infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Mycobacterium tuberculosis</i> H37Rv infection and associated clinical conditions	A
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GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	nicT	Mycobacterium bovis subsp. bovis AF2122/97	GAM353678 is a human miRNA-like oligonucleotide, which targets POSSIBLE NICKEL-TRANSPORT INTEGRAL MEMBRANE PROTEIN NICT (nicT, NC_002945 from 3123200 to 3124318 (+)), a bacterial target gene encoded by the <i>Mycobacterium bovis</i> subsp. bovis AF2122/97 genome, as part of an anti-bacterial host defense mechanism. nicT BINDING SITE 1 and nicT BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the nicT gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of nicT BINDING SITE 1 and nicT BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit nicT, a GAM353678 bacterial target gene which is associated	A
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					with Mycobacterium b ovis subsp bovis AF2122/97 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM 353678 include the diagnosis, prevention and treatment of Mycobacterium bovis subsp bovis AF2122/97 infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	nupC	Shigella flexneri 2a str. 301	GAM353678 is a human miRNA-like oligonucleotide, which targets permease of transport system for 3 nucleosides (nupC, NC_004337 from 2515842 to 2517083 (+)), a bacterial target gene encoded by the Shigella flexneri 2a str. 301 genome, as part of an anti-bacterial host defense mechanism. nupC BINDING SITE 1 through nupC BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the nupC gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of nupC BINDING SITE 1 through nupC BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit nupC, a GAM353678 bacterial target gene which is associated with Shigella flexneri 2a str. 301 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Shigella flexneri 2a str. 301 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	nupC	Escherichia coli CFT073	GAM353678 is a human miRNA-like oligonucleotide, which targets Nucleoside permease nupC (nupC, NC_004431 from 2795390 to 2796631 (+)), a bacterial target gene encoded by the Escherichia coli CFT073 genome, as part of an anti-bacterial host defense mechanism. nupC BINDING SITE 1 through nupC BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the nupC gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of nupC BINDING SITE 1 through nupC BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit nupC, a GAM353678 bacterial target gene which is associated with Escherichia coli CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT073 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	nupC	Shigella flexneri 2a str. 2	GAM353678 is a human miRNA-like oligonucleotide, which targets permease of transport system for 3 nucleosides	A

457T					(nupC, NC_0047 41 from 2494019 to 2495221 (+)), a bacterial target gene encoded by the Shigella flexneri 2a str. 2457T genome, as part of an anti-bacterial host defense mechanism. nupC BINDING SITE 1 through nupC BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the nupC gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of nupC BINDING SITE 1 through nupC BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit nupC, a GAM353678 bacterial target gene which is associated with Shigella flexneri 2a str. 2457T infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Shigella flexneri 2a str. 2457T infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	ompG	Escherichia coli CFT073	GAM353678 is a human miRNA-like oligonucleotide, which targets Outer membrane protein G precursor (ompG, NC_004431 from 16 24577 to 1625533 (+)), a bacterial target gene encoded by the Escherichia coli CFT073 genome, as part of an anti-bacterial host defense mechanism. ompG BINDING SITE 1 and ompG BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the ompG gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of ompG BINDING SITE 1 and ompG BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ompG, a GAM353678 bacterial target gene which is associated with Escherichia coli CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT073 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	orn	Yersinia pestis	GAM353678 is a human miRNA-like oligonucleotide, which targets oligoribonuclease (orn, NC_003143 from 378331 to 378876 (+)), a bacterial target gene encoded by the Yersinia pestis genome, as part of an anti-bacterial host defense mechanism. orn BINDING SITE is a bacterial target binding site that is found in the the 3' untranslated region of mRNA encoded by the orn gene, corresponding to a target binding site such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of orn BINDING SITE,	A

					and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit orn, a GAM353678 bacterial target gene which is associated with Yersinia pestis infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Yersinia pestis infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	oxyR	Salmonella enterica enterica serovar Typhi	GAM353678 is a human miRNA-like oligonucleotide, which targets hydrogen peroxide-inducible regulon activator (oxyR, NC_003198 from 3607204 to 3608121 (-)), a bacterial target gene encoded by the Salmonella enterica enterica serovar Typhi genome, as part of an anti-bacterial host defense mechanism. oxyR BINDING SITE 1 and oxyR BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the oxyR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of oxyR BINDING SITE 1 and oxyR BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit oxyR, a GAM353678 bacterial target gene which is associated with Salmonella enterica enterica serovar Typhi infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Salmonella enterica enterica serovar Typhi infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	oxyR	Salmonella typhimurium LT2	GAM353678 is a human miRNA-like oligonucleotide, which targets oxidative stress regulatory protein (oxyR, NC_003197 from 4343080 to 4343997 (+)), a bacterial target gene encoded by the Salmonella typhimurium LT2 genome, as part of an anti-bacterial host defense mechanism. oxyR BINDING SITE 1 and oxyR BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the oxyR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of oxyR BINDING SITE 1 and oxyR BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit oxyR, a GAM353678 bacterial target gene which is associated with Salmonella typhimurium LT2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Salmonella typhimurium LT2 infection and	A

					associated clinical conditions	
GAM35 3678	CAGCAGCA CACTGTGG TTTGTA	Human	oxyR	Salmonella enterica enterica serovar Typhi Ty2	GAM353678 is a human miRNA-like oligonucleotide, which targets hydrogen peroxide-inducible regulon activator (oxyR, NC_004631 from 3592864 to 3593781 (-)), a bacterial target gene encoded by the <i>Salmonella enterica enterica</i> serovar Typhi Ty2 genome, as part of an anti-bacterial host defense mechanism. oxyR BINDING SITE 1 and oxyR BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the oxyR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of oxyR BINDING SITE 1 and oxyR BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit oxyR, a bacterial target gene which is associated with <i>Salmonella enterica enterica</i> serovar Typhi Ty2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella enterica enterica</i> serovar Typhi Ty2 infection and associated clinical conditions	A
GAM35 3678	CAGCAGCA CACTGTGG TTTGTA	Human	pbpG	<i>Pseudomonas putida</i> KT2440	GAM353678 is a human miRNA-like oligonucleotide, which targets D-alanyl-D-alanine-endopeptidase (pbpG, NC_002947 from 4323707 to 4324633 (+)), a bacterial target gene encoded by the <i>Pseudomonas putida</i> KT2440 genome, as part of an anti-bacterial host defense mechanism. pbpG BINDING SITE 1 and pbpG BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the pbpG gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of pbpG BINDING SITE 1 and pbpG BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pbpG, a bacterial target gene which is associated with <i>Pseudomonas putida</i> KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Pseudomonas putida</i> KT2440 infection and associated clinical conditions	A
GAM35 3678	CAGCAGCA CACTGTGG TTTGTA	Human	pchA	<i>Pseudomonas aeruginosa</i> PA01	GAM353678 is a human miRNA-like oligonucleotide, which targets salicylate biosynthesis isochorismate synthase (pchA, NC_002516 from 4745120 to 4746550 (+)), a bacterial target gene encoded by the <i>Pseudomonas aeruginosa</i> PA01 genome, as part of an anti-bacterial host defense mechanism. pchA BINDING SITE 1 and pchA BINDING SITE 2 are bacterial target	A

binding sites that are found in the untranslated regions of mRNA encoded by the pchA gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of pchA BINDING SITE 1 and pchA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pchA, a GAM353678 bacterial target gene which is associated with *Pseudomonas aeruginosa* PA01 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Pseudomonas aeruginosa* PA01 infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	pcnA	Mycobacterium leprae	GAM353678 is a human miRNA-like oligonucleotide, which targets pcnA (pcnA, NC_002677 from 3248268 to 3249728 (-)), a bacterial target gene encoded by the <i>Mycobacterium leprae</i> genome, as part of an anti-bacterial host defense mechanism. pcnA BINDING SITE 1 and pcnA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the pcnA gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of pcnA BINDING SITE 1 and pcnA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pcnA, a GAM353678 bacterial target gene which is associated with <i>Mycobacterium leprae</i> infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Mycobacterium leprae</i> infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	phnV	Salmonella enterica enterica serovar Typhi	GAM353678 is a human miRNA-like oligonucleotide, which targets probable membrane component of 2-aminoethylphosphonate transporter (phnV, NC_003198 from 471575 to 472366 (-)), a bacterial target gene encoded by the <i>Salmonella enterica enterica</i> serovar Typhi genome, as part of an anti-bacterial host defense mechanism. phnV BINDING SITE 1 and phnV BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the phnV gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of phnV BINDING SITE 1 and phnV BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit phnV, a GAM353678	A

				bacterial target gene which is associated with <i>Salmonella enterica</i> serovar Typhi infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella enterica</i> serovar Typhi infection and associated clinical conditions		
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	phnV	<i>Salmonella enterica</i> serovar Typhi Ty2	GAM353678 is a human miRNA-like oligonucleotide, which targets probable membrane component of 2-aminoethylphosphonate transporter (phnV, NC_004631 from 2508735 to 2509526 (+)), a bacterial target gene encoded by the <i>Salmonella enterica</i> serovar Typhi Ty2 genome, as part of an anti-bacterial host defense mechanism. phnV BINDING SITE 1 and phnV BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the phnV gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of phnV BINDING SITE 1 and phnV BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit phnV, a bacterial target gene which is associated with <i>Salmonella enterica</i> serovar Typhi Ty2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella enterica</i> serovar Typhi Ty2 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	phoY2	<i>Mycobacterium bovis</i> subsp. <i>bovis</i> AF2122/97	GAM353678 is a human miRNA-like oligonucleotide, which targets PROBABLE PHOSPHATE-TRANSPORT SYSTEM TRANSCRIPTIONAL REGULATOR Y PROTEIN PHOY2 (phoY2, NC_002945 from 914388 to 915029 (-)), a bacterial target gene encoded by the <i>Mycobacterium bovis</i> subsp. <i>bovis</i> AF2122/97 genome, as part of an anti-bacterial host defense mechanism. phoY2 BINDING SITE 1 and phoY2 BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the phoY2 gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of phoY2 BINDING SITE 1 and phoY2 BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit phoY2, a bacterial target gene which is associated with <i>Mycobacterium bovis</i> subsp. <i>bovis</i> AF2122/97 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Mycobacterium bovis</i> subsp. <i>bovis</i> AF2122/97 infection and associated	A

				clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	phoY2	Mycobacterium tuberculosis H37Rv	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets phoY2 (phoY2, NC_000962 from 913556 to 914197 (-)), a bacterial target gene encoded by the Mycobacterium tuberculosis H37Rv genome, as part of an anti-bacterial host defense mechanism. phoY2 BINDING SITE 1 and phoY2 BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the phoY2 gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of phoY2 BINDING SITE 1 and phoY2 BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit phoY2, a bacterial target gene which is associated with Mycobacterium tuberculosis H37Rv infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Mycobacterium tuberculosis H37Rv infection and associated clinical conditions</p>
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	pilT	Pseudomonas putida KT2440	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets type IV pili twitching motility protein PilT (pilT, NC_002947 from 5816934 to 5817944 (-)), a bacterial target gene encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. pilT BINDING SITE 1 through pilT BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the pilT gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of pilT BINDING SITE 1 through pilT BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pilT, a bacterial target gene which is associated with Pseudomonas putida KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas putida KT2440 infection and associated clinical conditions</p>
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	polA	Mycobacterium leprae	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets DNA polymerase I (polA, NC_002677 from 1648220 to 1650955 (-)), a bacterial target gene encoded by the Mycobacterium leprae genome, as part of an anti-bacterial host defense mechanism. polA BINDING SITE 1 and polA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the polA gene, corresponding to target binding sites such as BINDING SITE I,</p>

BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of polA BINDING SITE 1 and polA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit polA, a GAM353678 bacterial target gene which is associated with Mycobacterium leprae infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Mycobacterium leprae infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	prcA	Mycobacterium leprae	GAM353678 is a human miRNA-like oligonucleotide, which targets proteasome [alpha]-type subunit 1 (prcA, NC_002677 from 1576553 to 1577350 (+)), a bacterial target gene encoded by the Mycobacterium leprae genome, as part of an anti-bacterial host defense mechanism. prcA BINDING SITE 1 and prcA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the prcA gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of prcA BINDING SITE 1 and prcA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit prcA, a GAM353678 bacterial target gene which is associated with Mycobacterium leprae infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Mycobacterium leprae infection and associated clinical conditions	A
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GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	pta	Pseudomonas putida KT2440	GAM353678 is a human miRNA-like oligonucleotide, which targets phosphate acetyltransferase (pta, NC_002947 from 891625 to 893712 (-)), a bacterial target gene encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. pta BINDING SITE 1 and pta BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the pta gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of pta BINDING SITE 1 and pta BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pta, a GAM353678 bacterial target gene which is associated with Pseudomonas putida KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis,	A
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GAM353678 is a human miRNA-like oligonucleotide, which targets phosphate acetyltransferase (pta, NC_002947 from 891625 to 893712 (-)), a bacterial target gene encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. pta BINDING SITE 1 and pta BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the pta gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of pta BINDING SITE 1 and pta BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pta, a GAM353678 bacterial target gene which is associated with Pseudomonas putida KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis,

					prevention and treatment of <i>Pseudomonas putida</i> KT 2440 infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	ptsH	Salmonella enterica serovar Typhi	GAM353678 is a human miRNA-like oligonucleotide, which targets phosphocarrier protein HPr (ptsH, NC_003198 from 2505403 to 2505660 (+)), a bacterial target gene encoded by the <i>Salmonella enterica enterica</i> serovar Typhi genome, as part of an anti-bacterial host defense mechanism. ptsH BINDING SITE is a bacterial target binding site that is found in the the 3' untranslated region of mRNA encoded by the ptsH gene, corresponding to a target binding site such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of ptsH BINDING SITE, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ptsH, a GAM353678 bacterial target gene which is associated with <i>Salmonella enterica enterica</i> serovar Typhi infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella enterica enterica</i> serovar Typhi infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	rbsR	Shigella flexneri 2a str. 301	GAM353678 is a human miRNA-like oligonucleotide, which targets regulator for rbs operon (rbsR, NC_004337 from 3947708 to 3948700 (+)), a bacterial target gene encoded by the <i>Shigella flexneri</i> 2a str. 301 genome, as part of an anti-bacterial host defense mechanism. rbsR BINDING SITE 1 through rbsR BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the rbsR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of rbsR BINDING SITE 1 through rbsR BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit rbsR, a GAM353678 bacterial target gene which is associated with <i>Shigella flexneri</i> 2a str. 301 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Shigella flexneri</i> 2a str. 301 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	rbsR	Shigella flexneri 2a str. 2457T	GAM353678 is a human miRNA-like oligonucleotide, which targets regulator for rbs operon (rbsR, NC_004741 from 3824594 to 3825577 (-)), a bacterial target gene encoded by the <i>Shigella flexneri</i> 2a str. 2457T genome, as part of an anti-bacterial host defense mechanism. rbsR BINDING SITE 1 through rbsR BINDING SITE 3 are bacterial target binding sites	A

that are found in the untranslated regions of mRNA encoded by the rbsR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of rbsR BINDING SITE 1 through rbsR BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit rbsR, a GAM353678 bacterial target gene which is associated with *Shigella flexneri* 2a str. 2457T infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Shigella flexneri* 2a str. 2457T infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	rbsR	Escherichia coli CFT073	GAM353678 is a human miRNA-like oligonucleotide, which targets Ribose operon repressor (rbsR, NC_004431 from 4439260 to 4440252 (+)), a bacterial target gene encoded by the Escherichia coli CFT073 genome, as part of an anti-bacterial host defense mechanism. rbsR BINDING SITE 1 through rbsR BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the rbsR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of rbsR BINDING SITE 1 through rbsR BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit rbsR, a GAM353678 bacterial target gene which is associated with Escherichia coli CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT073 infection and associated clinical conditions	A
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GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	recG	Mycobacterium leprae	GAM353678 is a human miRNA-like oligonucleotide, which targets ATP-dependent DNA helicase (recG, NC_002677 from 2014723 to 2016954 (-)), a bacterial target gene encoded by the Mycobacterium leprae genome, as part of an anti-bacterial host defense mechanism. recG BINDING SITE 1 and recG BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the recG gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of recG BINDING SITE 1 and recG BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit recG, a GAM353678	A
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GAM353678 is a human miRNA-like oligonucleotide, which targets ATP-dependent DNA helicase (recG, NC_002677 from 2014723 to 2016954 (-)), a bacterial target gene encoded by the Mycobacterium leprae genome, as part of an anti-bacterial host defense mechanism. recG BINDING SITE 1 and recG BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the recG gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of recG BINDING SITE 1 and recG BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit recG, a GAM353678

					bacterial target gene which is associated with Mycobacterium leprae infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Mycobacterium leprae infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	relA	Mycobacterium bovis subsp bovis AF2122/97	GAM353678 is a human miRNA-like oligonucleotide, which targets PROBABLE GTP PYROPHOSPHOKINASE RELA (ATP:GTP 3'-PYROPHOSPHOTRANSFERASE) (PPGPP SYNTHETASE I) ((P)PPGPP SYNTHETASE) (GTP DIPHOSPHOKINASE) (relA, NC_002945 from 2875274 to 2877646 (-)), a bacterial target gene encoded by the Mycobacterium bovis subsp bovis AF2122/97 genome, as part of an anti-bacterial host defense mechanism. relA BINDING SITE 1 and relA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the relA gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of relA BINDING SITE 1 and relA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit relA, a GAM353678 bacterial target gene which is associated with Mycobacterium bovis subsp bovis AF2122/97 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM 353678 include the diagnosis, prevention and treatment of Mycobacterium bovis subsp bovis AF2122/97 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	relA	Mycobacterium tuberculosis H37Rv	GAM353678 is a human miRNA-like oligonucleotide, which targets relA (relA, NC_000962 from 2907824 to 2910196 (-)), a bacterial target gene encoded by the Mycobacterium tuberculosis H37Rv genome, as part of an anti-bacterial host defense mechanism. relA BINDING SITE 1 and relA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the relA gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of relA BINDING SITE 1 and relA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit relA, a GAM353678 bacterial target gene which is associated with Mycobacterium tuberculosis H37Rv infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Mycobacterium tuberculosis H37Rv infection and associated clinical conditions	A

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	risA	Bordetella pertussis	GAM353678 is a human miRNA-like oligonucleotide, which targets tresponse regulator protein (risA, NC_002929 from 3765257 to 3765991 (-)), a bacterial target gene encoded by the Bordetella pertussis genome, as part of an anti-bacterial host defense mechanism. risA BINDING SITE 1 and risA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the risA gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of risA BINDING SITE 1 and risA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit risA, a GAM353678 bacterial target gene which is associated with Bordetella pertussis infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Bordetella pertussis infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	rpsT	Pseudomonas putida KT2440	GAM353678 is a human miRNA-like oligonucleotide, which targets ribosomal protein S20 (rpsT, NC_002947 from 707068 to 707346 (-)), a bacterial target gene encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. rpsT BINDING SITE 1 and rpsT BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the rpsT gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of rpsT BINDING SITE 1 and rpsT BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit rpsT, a GAM353678 bacterial target gene which is associated with Pseudomonas putida KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas putida KT2440 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	ruvB	Yersinia pestis	GAM353678 is a human miRNA-like oligonucleotide, which targets Holliday junction DNA helicase (ruvB, NC_003143 from 2336449 to 2337453 (+)), a bacterial target gene encoded by the Yersinia pestis genome, as part of an anti-bacterial host defense mechanism. ruvB BINDING SITE 1 and ruvB BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the ruvB gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of ruvB	A

					<p>BINDING SITE 1 and ruvB BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ruvB, a GAM353678 bacterial target gene which is associated with <i>Yersinia pestis</i> infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Yersinia pestis</i> infection and associated clinical conditions</p>		
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	ruvB	<i>Yersinia pestis</i> KIM	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets Holliday junction helicase subunit A (ruvB, NC_004088 from 2482031 to 2483035 (-)), a bacterial target gene encoded by the <i>Yersinia pestis</i> KIM genome, as part of an anti-bacterial host defense mechanism. ruvB BINDING SITE 1 and ruvB BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the ruvB gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of ruvB BINDING SITE 1 and ruvB BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ruvB, a GAM353678 bacterial target gene which is associated with <i>Yersinia pestis</i> KIM infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Yersinia pestis</i> KIM infection and associated clinical conditions</p>	A	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	selB	<i>Pseudomonas putida</i> KT2440	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets selenocysteine-specific translation elongation factor (selB, NC_002947 from 582133 to 584055 (+)), a bacterial target gene encoded by the <i>Pseudomonas putida</i> KT2440 genome, as part of an anti-bacterial host defense mechanism. selB BINDING SITE is a bacterial target binding site that is found in the the 3' untranslated region of mRNA encoded by the selB gene, corresponding to a target binding site such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of selB BINDING SITE, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit selB, a GAM353678 bacterial target gene which is associated with <i>Pseudomonas putida</i> KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Pseudomonas putida</i> KT2440 infection and associated clinical</p>	A	

				conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	SERPI Human NH 1	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets a human Serine proteinase inhibitor clade H (heat shock prote in 47) member 1; (SERPINH1, Accession number: NM_001235) as pa rt of a host response mechanism associated with a Escherichia coli CFT073, Streptococcus pneumoniae R6, Streptococcus pneumo niae TIGR4, Streptococcus pyogenes M1 GAS, Streptococcus pyoge nes MGAS315, Streptococcus pyogenes MGAS8232 and Streptococcus pyogenes SSI-1 infections. SERPINH1 BINDING SITE 1 and SERPINH1 BINDING SITE 2 are human target binding sites that are found in the untranslated region s of mRNA encoded by the SERPINH1 gene, corresponding to targe t binding sites such as BINDING SITE I, BINDING SITE II or BIN DING SITE III of Fig. 1. Additionally, using the binding site prediction system of the present invention GAM353678-A binds t o sequences on orthologous UTR of rat (NM_017173). The nucleoti de sequences of SERPINH1 BINDING SITE 1 and SERPINH1 BINDING S ITE 2, and the complementary secondary structure to the nucleo tide sequence of GAM353678 RNA are set forth in Tables 6-7, he reb y incorporated herein. Another function of GAM353678 is to inhibit SERPINH1, a GAM353678 human target gene which encodes a heat shock protein and s erpin, that may function as a chaperone for procollagen in the ER. SERPINH1 is associated with Escherichia coli CFT073, Stre ptococcus pneumoniae R6, Streptococcus pneumoniae TIGR4, Strep tococcus pyogenes M1 GAS, Streptococcus pyogenes MGAS315, Stre ptococcus pyogenes MGAS8232 and Streptococcus pyogenes SSI-1 i nfections, and therefore GAM353678 is associated with the abov ementioned infections, as part of a host response mechanism. A ccordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT073, Streptoco ccus pneumoniae R6, Streptococcus pneumoniae TIGR4, Streptococ cus pyogenes M1 GAS, Streptococcus pyogenes MGAS315, Streptoco ccus pyogenes MGAS8232 and Streptococcus pyogenes SSI-1 infect ions and associated clinical conditions. The function of SERPINH1 and its association with various dise ases and clinical conditions has been established by previous studies, as described hereinabove with reference to GAM839.</p>	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	sitD Shigella flexneri 2a str. 3 01	<p>GAM353678 is a human miRNA-like oligonucleotide, which targets Iron transport protein, inner membrane component (sitD, NC_00 4337 from 1405360 to 1406217 (-)), a bacterial target gene encoded by the Shigella flexneri 2a str. 301 genome, as part of an anti-bacterial host defense mechanism. sitD BINDING SITE 1 and sitD BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the sitD gene, corresponding to target</p>	A

binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of sitD BINDING SITE 1 and sitD BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit sitD, a GAM353678 bacterial target gene which is associated with *Shigella flexneri* 2a str. 301 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Shigella flexneri* 2a str. 301 infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	sitD	Shigella flexneri 2a str. 2457T	GAM353678 is a human miRNA-like oligonucleotide, which targets Iron transport protein, inner membrane component (sitD, NC_004741 from 1904666 to 1905523 (+)), a bacterial target gene encoded by the <i>Shigella flexneri</i> 2a str. 2457T genome, as part of an anti-bacterial host defense mechanism. sitD BINDING SITE 1 and sitD BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the sitD gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of sitD BINDING SITE 1 and sitD BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit sitD, a GAM353678 bacterial target gene which is associated with <i>Shigella flexneri</i> 2a str. 2457T infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Shigella flexneri</i> 2a str. 2457T infection and associated clinical conditions	A
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GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	speD	Salmonella enterica enterica serovar Typhi Ty2	GAM353678 is a human miRNA-like oligonucleotide, which targets S-adenosylmethionine decarboxylase proenzyme (speD, NC_004631 from 196380 to 197174 (-)), a bacterial target gene encoded by the <i>Salmonella enterica enterica</i> serovar Typhi Ty2 genome, as part of an anti-bacterial host defense mechanism. speD BINDING SITE 1 and speD BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the speD gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of speD BINDING SITE 1 and speD BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit speD, a GAM353678 bacterial target gene which is associated with <i>Salmonella enterica enterica</i> serovar Typhi Ty2 infection, as part of an anti-	A
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					bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella enterica enterica</i> serovar Typhi Ty2 infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	speD	<i>Salmonella enterica enterica</i> serovar Typhi	GAM353678 is a human miRNA-like oligonucleotide, which targets <i>S</i> -adenosylmethionine decarboxylase proenzyme (speD, NC_003198 from 196389 to 197183 (-)), a bacterial target gene encoded by the <i>Salmonella enterica enterica</i> serovar Typhi genome, as part of an anti-bacterial host defense mechanism. speD BINDING SITE 1 and speD BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the speD gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of speD BINDING SITE 1 and speD BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit speD, a bacterial target gene which is associated with <i>Salmonella enterica enterica</i> serovar Typhi infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella enterica enterica</i> serovar Typhi infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	speD	<i>Salmonella typhimurium</i> LT2	GAM353678 is a human miRNA-like oligonucleotide, which targets <i>S</i> -adenosylmethionine decarboxylase (speD, NC_003197 from 194201 to 194995 (-)), a bacterial target gene encoded by the <i>Salmonella typhimurium</i> LT2 genome, as part of an anti-bacterial host defense mechanism. speD BINDING SITE 1 and speD BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the speD gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of speD BINDING SITE 1 and speD BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit speD, a bacterial target gene which is associated with <i>Salmonella typhimurium</i> LT2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Salmonella typhimurium</i> LT2 infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	ssb	<i>Pseudomonas putida</i> KT2440	GAM353678 is a human miRNA-like oligonucleotide, which targets single-stranded DNA-binding protein (ssb, NC_002947 from 571027 to 571572 (+)), a bacterial target gene encoded by the	A

Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. *ssb* BINDING SITE 1 and *ssb* BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the *ssb* gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE I II of Fig. 1. The nucleotide sequences of *ssb* BINDING SITE 1 and *ssb* BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit *ssb*, a bacterial target gene which is associated with *Pseudomonas putida* KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Pseudomonas putida* KT 2440 infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGT	Human	<i>sseB</i>	<i>Escherichia coli</i> CFT073	GAM353678 is a human miRNA-like oligonucleotide, which targets Protein <i>sseB</i> (<i>sseB</i> , NC_004431 from 2922456 to 2923241 (-)), a bacterial target gene encoded by the <i>Escherichia coli</i> CFT 073 genome, as part of an anti-bacterial host defense mechanism. <i>sseB</i> BINDING SITE 1 and <i>sseB</i> BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the <i>sseB</i> gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of <i>sseB</i> BINDING SITE 1 and <i>sseB</i> BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit <i>sseB</i> , a bacterial target gene which is associated with <i>Escherichia coli</i> CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Escherichia coli</i> CFT073 infection and associated clinical conditions	A
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GAM353678	CAGCAGCA CACTGTGG TTTGT	Human	<i>tcfA</i>	<i>Bordetella pertussis</i>	GAM353678 is a human miRNA-like oligonucleotide, which targets tracheal colonization factor precursor (<i>tcfA</i> , NC_002929 from 1264436 to 1266379 (+)), a bacterial target gene encoded by the <i>Bordetella pertussis</i> genome, as part of an anti-bacterial host defense mechanism. <i>tcfA</i> BINDING SITE 1 and <i>tcfA</i> BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the <i>tcfA</i> gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of <i>tcfA</i> BINDING SITE 1 and <i>tcfA</i> BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678	A
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				RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit <i>tcfA</i> , a GAM353678 bacterial target gene which is associated with <i>Bordetella pertussis</i> infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Bordetella pertussis</i> infection and associated clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	<i>truA</i> <i>Mycobacterium leprae</i>	GAM353678 is a human miRNA-like oligonucleotide, which targets pseudouridylate synthase (<i>truA</i> , NC_002677 from 234 3329 to 2344078 (-)), a bacterial target gene encoded by the <i>Mycobacterium leprae</i> genome, as part of an anti-bacterial host defense mechanism. <i>truA</i> BINDING SITE 1 and <i>truA</i> BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the <i>truA</i> gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of <i>truA</i> BINDING SITE 1 and <i>truA</i> BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit <i>truA</i> , a GAM353678 bacterial target gene which is associated with <i>Mycobacterium leprae</i> infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Mycobacterium leprae</i> infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	truncated <i>Staphylococcus aureus</i> subsp. <i>aureus</i> MW2	GAM353678 is a human miRNA-like oligonucleotide, which targets truncated FmtB protein (truncated <i>fmtB</i> , NC_003923 from 2238 083 to 2240143 (-)), a bacterial target gene encoded by the <i>Staphylococcus aureus</i> subsp. <i>aureus</i> MW2 genome, as part of an anti-bacterial host defense mechanism. truncated <i>fmtB</i> BINDING SITE 1 and truncated <i>fmtB</i> BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the truncated <i>fmtB</i> gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of truncated <i>fmtB</i> BINDING SITE 1 and truncated <i>fmtB</i> BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit truncated <i>fmtB</i> , a GAM353678 bacterial target gene which is associated with <i>Staphylococcus aureus</i> subsp. <i>aureus</i> MW2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of <i>Staphylococcus aureus</i> subsp. <i>aureus</i> MW2 infection and associated	A

					clinical conditions	
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	uhpA	Yersinia pestis	GAM353678 is a human miRNA-like oligonucleotide, which targets two-component system response regulator (uhpA, NC_003143 from 4522790 to 4523380 (-)), a bacterial target gene encoded by the Yersinia pestis genome, as part of an anti-bacterial host defense mechanism. uhpA BINDING SITE 1 and uhpA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the uhpA gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of uhpA BINDING SITE 1 and uhpA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit uhpA, a bacterial target gene which is associated with Yersinia pestis infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Yersinia pestis infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	ung	Haemophilus influenzae Rd	GAM353678 is a human miRNA-like oligonucleotide, which targets uracil DNA glycosylase (ung, NC_000907 from 18676 to 19335 (+)), a bacterial target gene encoded by the Haemophilus influenzae Rd genome, as part of an anti-bacterial host defense mechanism. ung BINDING SITE 1 and ung BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the ung gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of ung BINDING SITE 1 and ung BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ung, a bacterial target gene which is associated with Haemophilus influenzae Rd infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Haemophilus influenzae Rd infection and associated clinical conditions	A
GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	vanB	Pseudomonas aeruginosa PA01	GAM353678 is a human miRNA-like oligonucleotide, which targets vanillate O-demethylase oxidoreductase (vanB, NC_002516 from 5504120 to 5505073 (+)), a bacterial target gene encoded by the Pseudomonas aeruginosa PA01 genome, as part of an anti-bacterial host defense mechanism. vanB BINDING SITE 1 and vanB BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the vanB gene, corresponding to target binding sites such as BINDING SITE I,	A

BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of vanB BINDING SITE 1 and vanB BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit vanB, a GAM353678 bacterial target gene which is associated with *Pseudomonas aeruginosa* PA01 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of *Pseudomonas aeruginosa* PA01 infection and associated clinical conditions

GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	yabO	Escherichia coli CFT073	GAM353678 is a human miRNA-like oligonucleotide, which targets Ribosomal large subunit pseudouridine synthase A (yabO, NC_004431 from 61489 to 62148 (-)), a bacterial target gene encoded by the Escherichia coli CFT073 genome, as part of an anti-bacterial host defense mechanism. yabO BINDING SITE 1 and yabO BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the yabO gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of yabO BINDING SITE 1 and yabO BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit yabO, a GAM353678 bacterial target gene which is associated with Escherichia coli CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT073 infection and associated clinical conditions	A
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GAM353678	CAGCAGCA CACTGTGG TTTGTA	Human	yciE	Escherichia coli CFT073	GAM353678 is a human miRNA-like oligonucleotide, which targets Protein yciE (yciE, NC_004431 from 1558641 to 1559147 (-)), a bacterial target gene encoded by the Escherichia coli CFT073 genome, as part of an anti-bacterial host defense mechanism. yciE BINDING SITE 1 and yciE BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the yciE gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of yciE BINDING SITE 1 and yciE BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit yciE, a GAM353678 bacterial target gene which is associated with Escherichia coli CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis,	A
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GAM353678 is a human miRNA-like oligonucleotide, which targets Protein yciE (yciE, NC_004431 from 1558641 to 1559147 (-)), a bacterial target gene encoded by the Escherichia coli CFT073 genome, as part of an anti-bacterial host defense mechanism. yciE BINDING SITE 1 and yciE BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the yciE gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of yciE BINDING SITE 1 and yciE BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit yciE, a GAM353678 bacterial target gene which is associated with Escherichia coli CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis,

prevention and treatment of Escherichia
coli CFT07 3 infection and associated
clinical conditions

Replace paragraph 0160 with the following paragraph.

Studies documenting the well known correlations between each of a plurality of GAM TARGET GENES that are described by Fig.1 and the known gene functions and related diseases are listed in Table 9, hereby incorporated herein. Specifically, in Table 9, lines 6046-6059 describes references of GAM target genes, as set forth in SEQ ID NO:348 in Table 8.

After paragraph 0160, add the following Table 9, paragraph, Table 11, paragraph, Table 12, paragraph, and Table 13.

Table 9:

TARGET	TARGET ORGANISM	REFERENCES
=====	=====	=====
MGAT5	Human	Demetriou, M.; Granovsky, M.; Quaggin, S.; Dennis, J. W.: Negative regulation of T-cell activation and autoimmunity by Mgat5 N-glycosylation. Nature 409: 733-739, 2001.
MGAT5	Human	Granovsky, M.; Fata, J.; Pawling, J.; Muller, W. J.; Khokha, R.; Dennis, J. W.: Suppression of tumor growth and metastasis in Mgat5-deficient mice. Nature Med.6: 306-12, 2000.
MGAT5	Human	Saito, H.; Nishikawa, A.; Gu, J.; Ihara, Y.; Soejima, H.; Wada, Y.; Sekiya, C.; Niikawa, N.; Taniguchi, N.: cDNA cloning and chromosomal mapping of human N-acetyl glucosaminyltransferase V+. Biochem. Biophys. Res. Commun. 198: 318-327, 1994.

Table 11, lines 275482-275565, shows data of GAM RNA SEQ ID NO:348 printed on microarray chip probes, as described in detail in Fig.17.

Table 11

PROBE SEQUENCE	PROBE TYPE	GAM RNA SEQ ID/ MIR NAME	GAM RNA/MIR SEQUENCE	LIB RARY	SIG NAL	BACKG ROUND Z-SCO RE	MISM ATCH Z-SCO RE
=====	=====	=====	=====	=====	=====	=====	=====
CCCAGCAGCAC ACTGTGGTTTG TACGCGATCCG TTATCGTTCGG TATCGAACGTA ACGAT	Predicted	348	CAGCAGCACACTGTGGTTTGTA	A2	638	4.2	3.2
CCCAGCAGCAC	Predicted	348	CAGCAGCACACTGTGGTTTGTA	D2	9435	16.6	20.9

ACTGTGGTTTG
TACGCGATCCG
TTATCGTTCGG
TATCGAACGTA
ACGAT

CCCAGCAGCAC Predicted 348 CAGCAGCAGCTGTGGTTTGTA E1 25910 14.8 27.5

ACTGTGGTTTG
TACGCGATCCG
TTATCGTTCGG
TATCGAACGTA
ACGAT

CCCAGCAGCAC Predicted 348 CAGCAGCAGCTGTGGTTTGTA F1 65518 12.0 30.2

ACTGTGGTTTG
TACGCGATCCG
TTATCGTTCGG
TATCGAACGTA
ACGAT

CCCAGCAGCAC Predicted 348 CAGCAGCAGCTGTGGTTTGTA G1 65518 10.1 29.3

ACTGTGGTTTG
TACGCGATCCG
TTATCGTTCGG
TATCGAACGTA
ACGAT

CCCAGCAGCAC Predicted 348 CAGCAGCAGCTGTGGTTTGTA H1 37067 9.9 28.2

ACTGTGGTTTG
TACGCGATCCG
TTATCGTTCGG
TATCGAACGTA
ACGAT

CCCAGCAGCAC Predicted 348 CAGCAGCAGCTGTGGTTTGTA A2 606 3.7 3.2

ACTGTGGTTTG
TACGGATCGTT
ATAACGATCCG
GTATCGAACGT
AACGA

CCCAGCAGCAC Predicted 348 CAGCAGCAGCTGTGGTTTGTA D2 7549 15.4 19.5

ACTGTGGTTTG
TACGGATCGTT
ATAACGATCCG
GTATCGAACGT
AACGA

CCCAGCAGCAC Predicted 348 CAGCAGCAGCTGTGGTTTGTA E1 20239 13.8 25.3

ACTGTGGTTTG
TACGGATCGTT
ATAACGATCCG
GTATCGAACGT
AACGA

CCCAGCAGCAC Predicted 348 CAGCAGCAGCTGTGGTTTGTA F1 65518 12.0 29.3

ACTGTGGTTTG
TACGGATCGTT
ATAACGATCCG
GTATCGAACGT
AACGA

CCCAGCAGCAC Predicted 348 CAGCAGCAGCTGTGGTTTGTA G1 65518 10.1 28.0

ACTGTGGTTTG
TACGGATCGTT
ATAACGATCCG
GTATCGAACGT
AACGA

CCCAGCAGCAC Predicted 348 CAGCAGCAGCTGTGGTTTGTA H1 27597 9.2 25.8

ACTGTGGTTTG

TACGGATCGTT
ATAACGATCCG
GTATCGAACGT
AACGA

Table 12, line 177, shows data relating to GAM RNA SEQ ID NO:348 that were validated by means of Wet Laboratory.

Table 12

GAM RNA SEQUENCE	VALIDATION METHOD	SIGNAL	BACKGROUND Z-SCORE	MISMATCH Z-SCORE	GAM RNA SEQ-ID
CAGCAGCACACTGTGGTTTGTA	Chip strong	65518	16.623587	30.172779	348

Table 13, lines 3-42, 47-69, 84-121, 143-179, 187-207, 210-256, 264-478 shows sequence data of GAMs associated with different bacterial infections.

Table 13

ROW#	INFECTION NAME	SEQ ID NOS OF GAMs ASSOCIATED WITH INFECTION
2	Bordetella pertussis	1, 6, 10, 11, 12, 13, 16, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 33, 34, 37, 41, 42, 43, 44, 47, 48, 49, 50, 52, 53, 54, 55, 57, 58, 59, 60, 63, 65, 66, 67, 68, 69, 70, 71, 75, 76, 77, 79, 84, 86, 87, 88, 89, 91, 94, 96, 97, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115, 117, 119, 120, 121, 122, 123, 125, 126, 127, 130, 131, 132, 133, 137, 138, 139, 140, 141, 142, 145, 147, 149, 150, 151, 154, 155, 156, 157, 158, 160, 161, 162, 164, 165, 166, 167, 168, 170, 171, 172, 173, 174, 175, 176, 177, 179, 180, 181, 183, 184, 185, 188, 191, 195, 196, 197, 204, 205, 211, 212, 214, 215, 216, 219, 220, 222, 225, 228, 230, 231, 233, 237, 239, 241, 242, 243, 244, 250, 251, 253, 262, 264, 265, 266, 268, 271, 272, 274, 276, 277, 280, 281, 282, 284, 285, 287, 288, 289, 290, 293, 294, 296, 297, 299, 300, 301, 302, 304, 306, 308, 310, 312, 317, 318, 321, 322, 324, 326, 327, 329, 330, 332, 333, 334, 335, 336, 339, 340, 342, 343, 345, 348, 349, 350, 351, 352, 353, 355, 356, 357, 358, 360, 361, 362, 364, 365, 366, 367, 368, 369, 370, 371, 372, 373, 374, 375, 376, 378, 380, 381, 382, 383, 384, 385 and 49788-55666.
3	Brucella 1330 suis	1, 6, 10, 11, 12, 13, 14, 16, 18, 19, 21, 23, 27, 32, 35, 37, 39, 40, 42, 47, 48, 49, 50, 52, 53, 58, 62, 63, 65, 68, 70, 71, 77, 79, 80, 85, 86, 89, 90, 98, 102, 105, 107, 108, 109, 111, 112, 114, 115, 119, 120, 124, 125, 121, 122, 123, 126, 132, 138, 141, 142, 143, 150, 151, 152, 154, 155, 156, 157, 158, 160, 161, 162, 164, 166, 168, 171, 172, 173, 175, 176, 177, 180, 181, 183, 185, 186, 190, 195, 198, 199, 200, 201, 205, 207, 211, 212, 214, 215, 217, 218, 219, 220, 221, 222, 225, 229, 230, 231, 233, 236, 237, 240, 241, 243, 244, 250, 251, 256, 258, 263, 264, 265, 266, 270, 277, 279, 280, 281, 282, 285, 287, 289, 290, 293, 294, 295, 297, 300, 302, 303, 306, 308, 310, 312, 315, 318, 319, 320, 321, 330, 331, 333, 334, 335, 342, 343, 347, 348, 349, 353, 354, 356, 357, 360, 361, 364, 365, 366, 368, 369, 370, 371, 373, 374, 375, 377, 381, 382, 384 and 55667-60259.
4	Chlamydia Trachomatis	2, 3, 4, 6, 7, 8, 9, 10, 13, 14, 16, 18, 19, 20, 21, 22, 25, 26, 27, 30, 31, 32, 33, 36, 37, 38, 40, 45, 46, 47, 48, 49, 51, 52, 55, 62, 63, 64, 67, 73, 74, 75, 78, 81, 82, 84, 85, 86, 87, 88, 91, 94, 95, 98, 99, 104, 105, 106, 111, 113, 116, 122, 124, 126, 128, 132, 133, 136, 138, 146, 148, 149, 152, 154, 155, 156, 157, 160, 164, 166, 167, 177, 179, 180, 181, 187, 188, 190, 192, 194, 198, 199, 200, 205, 207, 208, 209, 210, 211, 213, 214, 217,

- 218, 222, 224, 225, 226, 229, 232, 233, 235, 236, 239, 241, 242, 243, 244, 245, 248, 251, 252, 253, 254, 256, 257, 259, 262, 264, 265, 269, 270, 271, 272, 273, 274, 278, 279, 287, 288, 289, 293, 295, 296, 297, 298, 299, 302, 303, 305, 306, 309, 311, 312, 316, 318, 319, 320, 322, 323, 324, 325, 326, 327, 328, 330, 332, 333, 335, 338, 340, 341, 343, 344, 345, 348, 349, 350, 353, 354, 356, 363, 373, 384 and 60260-67437.
- 6 Chlamydomorph 3, 5, 6, 8, 9, 10, 13, 17, 20, 21, 22, 23, 25, 27, 28, 31, 32, 33, 37, 39, 45, 46, 47, 48, 50, 52, 55, 62, 63, 64, 66, 67, 69, 73, 74, 82, 84, 85, 88, 89, 90, 91, 92, 95, 101, 102, 104, 105, 111, 114, 124, 148, 125, 126, 128, 143, 146, 152, 159, 160, 161, 164, 165, 166, 168, 175, 180, 181, 187, 176, 177, 178, 179, 189, 190, 192, 194, 201, 203, 205, 207, 208, 209, 212, 213, 214, 217, 218, 221, 223, 224, 227, 232, 233, 234, 236, 238, 239, 241, 242, 243, 244, 245, 247, 248, 252, 257, 258, 259, 260, 262, 263, 271, 272, 274, 275, 279, 281, 282, 283, 286, 289, 295, 297, 298, 299, 302, 305, 306, 309, 311, 312, 314, 319, 323, 324, 325, 326, 327, 330, 333, 338, 340, 343, 344, 345, 346, 348, 349, 350, 352, 353, 354, 356, 363, 377, 382, 383, 384 and 68148-75439.
- 7 Chlamydomorphila 3, 5, 6, 8, 9, 10, 17, 20, 21, 22, 23, 25, 27, 31, 32, 33, 37, 39, 45, 46, 47, 50, 52, 55, 62, 63, 64, 66, 67, 69, 73, 74, 82, 90, 92, 95, 101, 84, 85, 88, 89, 102, 104, 105, 111, 114, 125, 126, 128, 148, 152, 159, 160, 161, 143, 146, 164, 165, 166, 168, 175, 176, 177, 178, 179, 187, 189, 190, 192, 194, 201, 203, 205, 207, 208, 209, 212, 180, 181, 213, 214, 217, 218, 221, 223, 224, 227, 232, 233, 234, 236, 238, 239, 241, 242, 243, 244, 245, 247, 248, 252, 257, 259, 260, 262, 263, 271, 272, 274, 275, 279, 281, 282, 283, 286, 289, 295, 297, 298, 299, 302, 305, 306, 309, 311, 312, 314, 319, 323, 325, 326, 327, 330, 333, 338, 340, 343, 344, 345, 346, 348, 349, 350, 352, 353, 354, 356, 363, 377, 382, 383, 384 and 75440-82241.
- 10 Escherichia coli CFT 073 1, 2, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 16, 17, 18, 19, 21, 22, 23, 25, 26, 27, 28, 30, 31, 33, 34, 35, 36, 37, 39, 40, 42, 43, 45, 46, 47, 48, 49, 50, 51, 52, 53, 55, 56, 57, 58, 59, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 75, 76, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 99, 100, 101, 102, 103, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115, 116, 119, 120, 121, 122, 123, 124, 125, 126, 129, 131, 132, 133, 135, 136, 137, 138, 140, 141, 142, 143, 145, 146, 147, 148, 152, 154, 155, 156, 157, 158, 160, 161, 162, 163, 164, 165, 166, 167, 168, 171, 173, 174, 175, 176, 177, 179, 180, 181, 182, 184, 185, 186, 190, 191, 192, 193, 195, 196, 197, 203, 204, 205, 206, 207, 208, 209, 210, 211, 212, 213, 214, 215, 216, 217, 218, 220, 221, 222, 223, 224, 225, 229, 230, 231, 232, 233, 234, 235, 236, 237, 238, 239, 241, 242, 243, 244, 245, 246, 247, 248, 249, 250, 251, 252, 253, 254, 255, 256, 257, 258, 260, 261, 262, 265, 266, 267, 268, 270, 271, 272, 274, 276, 277, 278, 279, 280, 281, 282, 283, 284, 285, 287, 288, 289, 290, 291, 292, 293, 294, 295, 296, 297, 299, 300, 301, 302, 303, 305, 306, 307, 308, 309, 310, 311, 312, 314, 315, 316, 317, 318, 321, 322, 323, 324, 325, 326, 327, 329, 331, 332, 333, 334, 335, 336, 337, 338, 339, 340, 343, 344, 345, 347, 348, 349, 350, 351, 352, 353, 354, 355, 356, 357, 358, 359, 360, 361, 364, 365, 367, 368, 369, 370, 372, 373, 374, 375, 376, 378, 380, 381, 382, 383, 384, 385 and 90623-103607.
- 11 Haemophilus influenzae Rd 2, 3, 5, 6, 7, 8, 9, 10, 13, 15, 19, 20, 21, 22, 25, 26, 27, 30, 31, 32, 33, 34, 37, 38, 40, 41, 45, 46, 48, 49, 50, 51, 52, 66, 67, 68, 73, 53, 55, 62, 63, 64, 78, 81, 83, 84, 85, 88, 90, 91, 92, 98, 101, 105, 106, 111, 116, 117, 119, 122, 123, 124, 125, 126, 134, 138, 144, 146, 149, 151, 152, 155, 156, 160, 161, 164, 165, 166, 169, 171, 172, 174, 176, 177, 179, 180, 183, 190, 197, 198, 199, 200, 201, 203, 205, 207, 208, 211, 213, 214, 218, 221, 223, 226, 228, 229, 234, 236, 239, 240, 242, 244, 247, 248, 251, 254, 255, 256, 259, 262, 263, 264, 271, 272, 274, 277, 279, 281, 282, 283, 295, 296, 299, 302, 305, 306, 308, 311, 312, 313, 316, 317, 318, 319, 322, 323, 324, 325, 326, 327, 329, 333, 335, 338, 339, 340, 343, 344, 345, 348, 351, 353, 354, 356, 365,

- 368, 371, 375, 377, 379, 380, 385 and 103608-111433.
- 12 *Leptospira interrogans* serovar lai str. 56601 1, 3, 5, 7, 8, 10, 13, 19, 22, 25, 32, 38, 39, 41, 48, 49, 52, 67, 71, 73, 84, 85, 90, 91, 93, 95, 117, 124, 128, 164, 174, 192, 193, 203, 178, 179, 187, 190, 207, 225, 226, 227, 229, 238, 244, 258, 259, 262, 272, 279, 193, 203, 256, 257, 295, 298, 299, 303, 306, 307, 316, 324, 327, 333, 338, 340, 344, 348, 376, 379, 384 and 111434-116384.
- 15 *Mycobacterium bovis* AF2122/9 subsp *bovis* 7 1, 3, 4, 5, 6, 7, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 31, 32, 33, 36, 37, 39, 41, 42, 52, 43, 45, 46, 47, 48, 50, 51, 53, 54, 55, 57, 58, 59, 73, 75, 76, 60, 61, 62, 71, 64, 65, 66, 67, 68, 69, 70, 77, 78, 79, 80, 83, 84, 86, 87, 88, 89, 100, 101, 102, 90, 91, 93, 96, 97, 99, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115, 116, 117, 118, 119, 120, 121, 122, 123, 125, 127, 130, 131, 132, 133, 134, 135, 137, 138, 139, 141, 142, 143, 144, 145, 146, 147, 148, 149, 150, 151, 152, 154, 155, 156, 157, 158, 159, 160, 161, 162, 163, 165, 166, 167, 168, 170, 171, 172, 173, 174, 175, 176, 177, 178, 179, 180, 181, 183, 184, 185, 188, 189, 190, 191, 193, 194, 195, 196, 197, 198, 199, 200, 201, 202, 203, 204, 205, 206, 207, 208, 210, 211, 212, 213, 214, 215, 216, 218, 219, 220, 222, 225, 230, 231, 233, 236, 237, 239, 240, 241, 242, 243, 244, 245, 246, 250, 251, 252, 253, 254, 255, 256, 257, 261, 262, 263, 264, 265, 266, 267, 268, 270, 271, 273, 276, 277, 278, 280, 281, 282, 283, 285, 286, 287, 288, 289, 290, 291, 292, 293, 294, 296, 297, 299, 300, 302, 303, 304, 305, 306, 308, 310, 312, 313, 314, 315, 318, 320, 321, 322, 323, 324, 325, 326, 327, 329, 330, 331, 332, 333, 334, 335, 336, 337, 341, 342, 345, 346, 347, 348, 349, 350, 351, 352, 353, 355, 356, 357, 358, 360, 361, 362, 364, 365, 366, 367, 369, 370, 371, 372, 373, 374, 375, 376, 378, 380, 381, 382, 383, 384, 385 and 127919-137561.
- 16 *Mycobacterium leprae* 3, 4, 5, 6, 7, 12, 13, 14, 15, 18, 19, 21, 22, 23, 24, 26, 29, 31, 32, 33, 36, 37, 39, 41, 42, 43, 45, 46, 47, 48, 49, 50, 53, 54, 68, 69, 70, 57, 59, 62, 65, 71, 73, 74, 75, 76, 78, 81, 83, 84, 86, 90, 94, 96, 98, 101, 103, 106, 107, 109, 110, 111, 112, 113, 114, 115, 116, 118, 119, 120, 121, 123, 131, 133, 134, 135, 137, 142, 143, 144, 145, 146, 147, 149, 154, 156, 157, 158, 159, 161, 162, 163, 165, 166, 167, 171, 172, 173, 174, 175, 176, 179, 183, 184, 185, 187, 188, 189, 190, 193, 196, 197, 198, 199, 200, 201, 202, 204, 205, 206, 211, 212, 214, 215, 216, 218, 219, 220, 221, 223, 224, 225, 228, 230, 231, 232, 233, 234, 235, 236, 237, 241, 242, 243, 245, 249, 250, 251, 253, 254, 256, 258, 261, 263, 265, 267, 268, 269, 271, 274, 276, 277, 280, 281, 284, 288, 289, 290, 291, 293, 294, 295, 296, 297, 299, 300, 301, 302, 303, 305, 306, 307, 309, 310, 311, 312, 313, 314, 315, 318, 320, 321, 323, 324, 327, 329, 330, 331, 332, 333, 334, 335, 336, 337, 338, 339, 340, 341, 343, 345, 346, 347, 348, 349, 353, 355, 356, 357, 358, 360, 361, 364, 365, 368, 369, 370, 371, 372, 374, 375, 376, 377, 378, 380, 381, 382, 383 and 137562-144598.
- 18 *Mycobacterium tuberculosis* H37Rv 1, 3, 4, 5, 6, 7, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 31, 32, 33, 37, 39, 41, 42, 43, 50, 51, 52, 45, 46, 47, 48, 53, 54, 55, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 73, 75, 76, 77, 78, 79, 80, 83, 84, 86, 87, 88, 89, 90, 91, 93, 94, 96, 97, 99, 100, 101, 102, 103, 104, 105, 107, 108, 109, 110, 111, 112, 113, 114, 115, 116, 117, 118, 119, 120, 121, 122, 123, 125, 127, 130, 131, 132, 133, 134, 135, 137, 138, 139, 141, 142, 143, 144, 145, 146, 147, 148, 149, 150, 151, 152, 153, 154, 155, 156, 157, 158, 159, 160, 161, 162, 163, 165, 166, 167, 168, 170, 171, 172, 173, 174, 175, 176, 177, 178, 179, 180, 181, 183, 184, 185, 188, 189, 190, 191, 194, 195, 196, 197, 198, 199, 200, 201, 202, 203, 204, 205, 206, 207, 208, 210, 211, 212, 213, 214, 215, 216, 218, 219, 220, 222, 225, 230, 231, 233, 234, 236, 237, 239, 240, 241, 242, 243, 244, 245, 246, 250, 251, 252, 253, 254, 255, 256, 257, 261, 262, 263, 264, 265, 266, 267, 268, 270, 271, 272, 273, 274, 276, 277, 278, 280, 281, 282, 283, 284, 285, 286, 287, 288, 289, 290, 291, 292, 293, 294, 296, 297, 298, 299, 300, 302, 303, 304,

- 305, 306, 308, 310, 312, 313, 314, 315, 318, 320, 321, 323, 324, 325, 326, 327, 329, 330, 331, 332, 333, 334, 335, 336, 337, 341, 342, 345, 346, 347, 348, 349, 350, 351, 352, 353, 355, 356, 357, 358, 360, 361, 362, 364, 365, 366, 367, 368, 369, 370, 371, 372, 373, 374, 375, 376, 378, 380, 381, 382, 383, 384, 385 and 146807-155497.
- 20 *Neisseria meningitidis* Z2491 1, 6, 7, 8, 10, 12, 15, 17, 21, 22, 26, 28, 30, 37, 39, 40, 45, 49, 52, 56, 58, 60, 62, 63, 67, 70, 76, 86, 89, 90, 91, 96, 98, 102, 103, 105, 107, 108, 109, 111, 112, 113, 114, 115, 122, 123, 124, 125, 126, 127, 133, 138, 141, 142, 143, 145, 147, 148, 149, 152, 157, 158, 164, 165, 166, 170, 171, 175, 176, 178, 181, 183, 187, 189, 197, 203, 217, 218, 219, 220, 221, 222, 225, 229, 230, 231, 237, 239, 243, 245, 247, 248, 251, 253, 254, 256, 257, 258, 259, 264, 265, 268, 273, 281, 282, 283, 285, 287, 289, 290, 293, 294, 295, 297, 300, 302, 306, 308, 314, 315, 316, 319, 321, 322, 325, 327, 329, 332, 333, 334, 338, 340, 341, 344, 346, 348, 349, 350, 351, 354, 355, 356, 365, 371, 372, 375, 376, 380, 381, 382, 384 and 155834-160603.
- 21 *Pseudomonas aeruginosa* PA01 1, 2, 6, 10, 11, 12, 13, 14, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 33, 34, 35, 36, 37, 41, 42, 56, 43, 45, 46, 48, 49, 50, 52, 54, 55, 57, 58, 59, 60, 62, 63, 64, 76, 77, 78, 79, 81, 85, 86, 67, 68, 69, 70, 71, 73, 82, 83, 84, 86, 87, 88, 89, 90, 91, 94, 95, 96, 97, 99, 100, 101, 102, 103, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115, 116, 118, 119, 120, 121, 122, 123, 124, 125, 126, 127, 130, 131, 134, 137, 138, 139, 140, 141, 142, 144, 147, 149, 150, 151, 152, 154, 155, 156, 157, 158, 160, 161, 162, 163, 164, 165, 166, 167, 168, 169, 170, 171, 172, 173, 174, 175, 176, 177, 179, 180, 181, 183, 184, 185, 188, 190, 192, 193, 194, 195, 196, 197, 202, 204, 205, 208, 210, 211, 212, 213, 214, 215, 216, 218, 220, 222, 225, 228, 229, 230, 231, 232, 233, 236, 237, 241, 242, 243, 244, 250, 251, 253, 258, 262, 264, 265, 266, 267, 268, 270, 271, 272, 273, 274, 276, 277, 280, 281, 282, 283, 285, 286, 287, 288, 289, 290, 291, 292, 293, 294, 296, 297, 298, 299, 300, 301, 302, 306, 312, 314, 318, 319, 320, 321, 323, 324, 325, 327, 329, 330, 331, 333, 334, 335, 336, 339, 340, 341, 342, 343, 345, 347, 348, 349, 350, 351, 352, 353, 355, 356, 357, 358, 360, 361, 362, 364, 365, 366, 367, 368, 369, 370, 371, 372, 373, 374, 375, 376, 378, 380, 381, 382, 383, 384 and 160604-170274.
- 22 *Pseudomonas K putida* T2440 1, 5, 7, 9, 10, 11, 12, 13, 14, 16, 18, 19, 23, 24, 25, 26, 27, 28, 29, 31, 33, 34, 36, 37, 38, 39, 41, 42, 43, 44, 45, 46, 47, 49, 50, 51, 52, 53, 54, 55, 57, 58, 59, 61, 64, 65, 66, 68, 69, 70, 71, 73, 76, 84, 85, 86, 88, 89, 91, 94, 98, 99, 101, 102, 103, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115, 117, 118, 119, 120, 121, 122, 123, 125, 126, 131, 132, 133, 134, 135, 137, 138, 140, 141, 142, 143, 144, 145, 146, 147, 148, 149, 150, 151, 152, 153, 154, 155, 156, 157, 158, 159, 160, 161, 162, 163, 164, 166, 167, 168, 171, 172, 173, 174, 175, 176, 177, 179, 180, 181, 183, 184, 185, 187, 190, 191, 193, 195, 196, 197, 202, 204, 205, 207, 211, 212, 214, 215, 216, 220, 221, 222, 225, 228, 229, 230, 231, 232, 233, 234, 236, 237, 240, 241, 242, 243, 244, 248, 250, 251, 253, 255, 258, 264, 265, 266, 267, 270, 271, 272, 274, 276, 277, 280, 281, 282, 283, 284, 285, 287, 288, 289, 290, 291, 292, 293, 294, 295, 296, 297, 298, 299, 300, 301, 302, 303, 304, 305, 306, 308, 310, 312, 313, 314, 316, 317, 318, 320, 321, 322, 323, 324, 327, 329, 333, 334, 335, 336, 337, 342, 343, 345, 346, 347, 348, 349, 350, 351, 352, 353, 354, 355, 356, 357, 358, 360, 361, 364, 365, 366, 367, 368, 369, 370, 371, 373, 374, 375, 376, 377, 378, 380, 381, 382, 383, 384, 385 and 170275-178543.
- 24 *Salmonella enterica* serovar 1, 2, 3, 4, 6, 7, 8, 9, 10, 11, 12, 13, 16, 17, 18, 19, 20, 21, 22, 23, 25, 26, 27, 28, 30, 31, 32, 33, 35, 37, 38, 39, 51, 40, 42, 43, 45, 46, 47, 48, 49, 50, 52, 55, 56, 57, 58, 59, 60, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 75,

Typ hi	7, 79, 80, 81, 83, 84, 86, 88, 89, 90, 91, 92, 94, 95, 98, 99, 100, 101, 102, 105, 106, 107, 108, 109, 111, 112, 113, 114, 115, 116, 119, 120, 121, 122, 123, 124, 125, 126, 127, 129, 131, 132, 133, 135, 136, 137, 138, 142, 143, 144, 145, 146, 147, 148, 150, 152, 153, 154, 155, 156, 157, 158, 160, 161, 162, 163, 164, 165, 166, 167, 171, 172, 173, 174, 175, 176, 177, 179, 180, 181, 182, 183, 185, 187, 188, 189, 190, 191, 192, 193, 194, 195, 196, 197, 198, 199, 200, 201, 202, 203, 204, 205, 206, 208, 212, 213, 214, 215, 216, 217, 218, 219, 220, 221, 222, 223, 225, 226, 229, 230, 231, 232, 233, 234, 235, 236, 237, 238, 239, 240, 241, 242, 243, 244, 245, 247, 248, 250, 251, 252, 253, 254, 255, 256, 257, 260, 261, 262, 263, 265, 266, 269, 270, 271, 272, 274, 276, 277, 278, 280, 281, 282, 283, 284, 285, 287, 288, 289, 290, 291, 292, 293, 294, 295, 296, 297, 298, 299, 300, 301, 302, 303, 304, 305, 306, 308, 311, 312, 314, 315, 318, 319, 323, 324, 325, 327, 328, 329, 330, 331, 332, 333, 334, 335, 336, 337, 339, 340, 341, 342, 343, 344, 345, 346, 347, 348, 349, 351, 352, 353, 354, 355, 356, 357, 358, 360, 361, 364, 365, 366, 367, 369, 370, 371, 373, 374, 375, 376, 378, 379, 380, 381, 382, 383, 384, 385 and 179915-190940.
25 Salmonella enterica enterica serovar Typ hi Ty2	1, 2, 3, 4, 6, 7, 8, 9, 10, 11, 12, 13, 14, 16, 17, 18, 19, 20, 21, 22, 23, 25, 26, 27, 28, 30, 31, 32, 33, 35, 37, 38, 50, 39, 40, 42, 43, 45, 46, 47, 48, 49, 51, 52, 55, 56, 57, 58, 59, 69, 70, 71, 72, 73, 60, 62, 63, 64, 65, 66, 67, 68, 75, 77, 79, 80, 81, 83, 84, 85, 86, 88, 89, 90, 91, 94, 95, 98, 99, 100, 101, 102, 105, 106, 107, 108, 109, 111, 112, 113, 114, 115, 116, 119, 120, 121, 122, 123, 124, 125, 126, 127, 129, 131, 132, 133, 135, 136, 137, 138, 142, 143, 144, 145, 146, 147, 148, 150, 152, 153, 154, 155, 156, 157, 158, 160, 161, 162, 163, 164, 165, 166, 167, 171, 172, 173, 174, 175, 176, 177, 179, 180, 181, 182, 183, 185, 187, 188, 189, 190, 191, 192, 193, 194, 195, 196, 197, 198, 199, 200, 201, 202, 203, 204, 205, 206, 208, 212, 213, 214, 215, 216, 217, 218, 219, 220, 221, 222, 223, 225, 226, 229, 230, 231, 232, 233, 234, 235, 236, 237, 238, 239, 240, 241, 242, 243, 244, 245, 247, 248, 250, 251, 252, 253, 254, 255, 256, 257, 260, 261, 262, 263, 265, 266, 269, 270, 271, 272, 274, 276, 277, 278, 280, 281, 282, 283, 284, 285, 287, 288, 289, 290, 291, 292, 293, 294, 295, 296, 297, 298, 299, 300, 301, 302, 303, 304, 305, 306, 308, 311, 312, 314, 315, 318, 319, 323, 324, 325, 327, 328, 329, 330, 331, 332, 333, 334, 335, 336, 337, 338, 339, 340, 341, 342, 343, 344, 345, 346, 347, 348, 349, 351, 352, 353, 354, 355, 356, 357, 358, 360, 361, 364, 365, 366, 367, 369, 370, 371, 373, 374, 375, 376, 378, 379, 380, 381, 382, 383, 384, 385 and 190941-201927.
26 Salmonella typhimurium LT2	1, 2, 3, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 48, 49, 36, 37, 38, 39, 42, 43, 45, 46, 47, 50, 51, 52, 54, 55, 56, 57, 58, 59, 60, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 75, 77, 79, 82, 83, 84, 86, 88, 89, 90, 91, 94, 95, 96, 100, 101, 102, 103, 104, 105, 107, 108, 109, 111, 112, 113, 114, 115, 116, 119, 120, 121, 122, 123, 124, 125, 126, 127, 129, 131, 132, 133, 135, 137, 138, 142, 143, 144, 145, 146, 147, 148, 149, 150, 151, 152, 154, 155, 156, 157, 158, 160, 161, 162, 163, 164, 165, 166, 167, 168, 170, 171, 172, 173, 174, 175, 176, 177, 179, 180, 181, 182, 183, 185, 187, 188, 189, 190, 192, 193, 194, 195, 196, 197, 198, 199, 200, 201, 202, 203, 204, 205, 206, 207, 208, 211, 212, 213, 214, 215, 216, 217, 218, 219, 220, 221, 222, 225, 226, 227, 228, 229, 230, 231, 232, 233, 234, 235, 236, 237, 238, 239, 240, 241, 242, 243, 244, 245, 247, 248, 249, 250, 251, 252, 253, 255, 256, 257, 258, 260, 261, 262, 263, 266, 267, 268, 270, 271, 272, 273, 274, 275, 276, 279, 280, 281, 282, 283, 285, 287, 288, 289, 290, 291, 292, 293, 294, 296, 297, 298, 299, 300, 302, 303, 306, 307, 308, 309, 310, 311, 312, 314, 315, 317, 318, 319, 323, 324, 325, 326, 327, 328, 329, 330, 331, 332, 333, 334, 335, 336, 337, 338, 340, 341, 342, 343, 344, 345, 347, 348, 349, 350, 351, 352, 353, 354, 355, 356, 357, 358, 359, 360, 361, 362, 363, 364, 365, 366, 368, 369, 370, 371, 373, 374, 375, 376, 379, 380, 381,

382, 383, 384, 385 and 201928-215605.

- 27 *Shigella* 2a 1, 2, 5, 6, 9, 10, 11, 12, 13, 14, 16, 17, 18, 19, 21,
flexneri 22, 23, 24, 25, 26, 27, 28, 29, 30, 32, 33, 35, 36, 37, 50, 51,
 str.2457T 38, 39, 40, 41, 42, 43, 46, 47, 48, 49, 52, 54, 55, 56, 57, 58,
 59, 62, 63, 65, 66, 67, 68, 69, 70, 71, 73, 76, 78, 80,
 83, 84, 85, 86, 87, 88, 89, 90, 91, 93, 94, 95, 97, 99, 101,
 102, 103, 104, 105, 107, 108, 109, 110, 111, 112, 113, 114,
 115, 116, 117, 119, 120, 121, 122, 123, 124, 125, 126, 129,
 131, 132, 133, 134, 135, 136, 137, 138, 139, 141, 142, 143,
 145, 146, 147, 148, 149, 150, 151, 152, 154, 155, 156, 157,
 158, 160, 161, 162, 163, 164, 165, 166, 167, 171, 172, 173,
 174, 175, 176, 177, 179, 180, 181, 182, 184, 185, 187, 190,
 191, 195, 196, 197, 198, 199, 200, 201, 202, 203, 204, 205,
 207, 208, 212, 213, 214, 216, 218, 220, 221, 222, 223, 224,
 225, 229, 230, 231, 232, 233, 234, 236, 237, 238, 239, 240,
 241, 242, 243, 244, 245, 247, 248, 250, 251, 252, 253, 254,
 255, 256, 257, 260, 261, 262, 263, 265, 268, 270, 271, 272,
 274, 276, 277, 278, 279, 280, 281, 282, 283, 284, 285, 287,
 288, 289, 290, 291, 292, 293, 295, 296, 297, 298, 299, 300,
 301, 302, 304, 306, 307, 308, 309, 310, 311, 312, 314, 315,
 316, 317, 318, 320, 321, 322, 323, 324, 325, 327, 328, 329,
 331, 332, 333, 334, 335, 336, 337, 338, 339, 340, 341, 343,
 344, 345, 346, 347, 348, 349, 350, 351, 352, 353, 354, 356,
 357, 358, 359, 360, 361, 362, 364, 365, 366, 367, 368, 369,
 371, 373, 374, 375, 376, 379, 380, 381, 382, 383, 384, 385
 and 215606-226197.
- 28 *Shigella* 2a 1, 2, 5, 6, 9, 10, 11, 12, 13, 14, 16, 17, 18, 19, 21,
flexneri 22, 23, 24, 25, 26, 27, 28, 29, 30, 32, 33, 35, 36, 37, 52,
 str. 301 39, 40, 41, 42, 43, 46, 47, 48, 49, 50, 51, 54, 55, 56, 57,
 58, 59, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 73, 76, 77, 78,
 80, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 97,
 99, 101, 102, 103, 104, 105, 107, 108, 109, 110, 111, 112,
 113, 114, 115, 116, 119, 120, 121, 122, 123, 124, 125,
 126, 129, 132, 133, 134, 135, 136, 137, 138, 141, 142,
 143, 144, 145, 146, 147, 148, 149, 150, 151, 152, 154, 155,
 156, 157, 158, 159, 160, 161, 162, 163, 164, 165, 166, 167,
 168, 171, 172, 173, 174, 175, 176, 177, 179, 180,
 181, 182, 184, 185, 187, 190, 191, 195, 196, 197, 198, 199,
 200, 201, 202, 203, 205, 207, 208, 210, 212, 213, 214, 216,
 217, 218, 220, 221, 222, 223, 224, 225, 229, 230, 231,
 232, 233, 234, 236, 237, 238, 239, 240, 241, 242, 243,
 244, 245, 247, 248, 250, 251, 252, 253, 254, 255, 256, 257,
 260, 262, 263, 264, 265, 266, 268, 269, 270, 271, 272, 274,
 276, 277, 278, 279, 280, 281, 282, 283, 284, 285, 287, 288,
 289, 290, 291, 292, 293, 295, 296, 297, 298, 299, 300, 301,
 302, 304, 306, 308, 309, 311, 312, 314, 315, 316, 317, 318,
 320, 321, 323, 324, 325, 327, 328, 329, 331, 333, 334, 335,
 336, 337, 338, 339, 340, 341, 343, 344, 345, 346, 347, 348,
 349, 350, 351, 352, 353, 354, 356, 357, 358, 359, 360, 361,
 362, 364, 365, 366, 367, 368, 369, 371, 373, 374, 375, 376,
 378, 379, 380, 381, 382, 383, 384, 385 and 226198-237003.
- 29 *Staphyl* 2, 5, 7, 8, 9, 10, 13, 16, 19, 22, 25, 27, 31,
Ococcus 32, 33, 35, 36, 38, 39, 40, 41, 45, 46, 47
Aureu , 48, 50, 51, 52, 55, 62, 63, 67, 71, 73, 81, 83, 84, 85,
s subsp 90, 91, 92, 93, 95, 98, 100, 101, 105, 106, 111, 113, 116,
. aureus 119, 120, 124, 131, 133, 138, 139, 146, 147, 149, 152
 Mu50 , 153, 156, 160, 161, 162, 165, 166, 169, 171, 172, 174,
 177, 179, 180, 181, 190, 192, 203, 204, 205, 207, 208, 213,
 214, 217, 218, 222, 228, 231, 232, 236, 238, 240, 242, 244,
 245, 247, 248, 252, 254, 256, 259, 261, 262, 270, 271, 272,
 274, 275, 287, 293, 294, 299, 301, 302, 305, 306, 308, 309,
 311, 316, 317, 323, 324, 325, 326, 327, 332, 333, 334, 335,
 337, 339, 340, 342, 343, 344, 345, 346, 348, 349, 351, 353,
 354, 356, 363, 365, 368, 371, 375, 379, 381 and 237004-244310.
- 30 *Staphyl* 2, 5, 7, 8, 10, 13, 16, 19, 22, 25, 27, 30, 31, 32,
ococuss 33, 38, 39, 40, 41, 45, 46, 47, 48, 50, 51, 52, 55,

- subsp. aureus MW2 62, 63, 67, 71, 72, 73, 78, 81, 83, 84, 90, 91, 92, 93, 95, 98, 100, 101, 105, 106, 109, 111, 113, 117, 119, 120, 124, 126, 128, 130, 131, 133, 134, 138, 139, 143, 149, 152, 153, 156, 160, 161, 162, 166, 169, 171, 172, 174, 177, 179, 180, 181, 182, 190, 192, 203, 204, 205, 207, 208, 213, 214, 217, 218, 222, 228, 231, 232, 236, 238, 242, 244, 247, 248, 252, 254, 256, 257, 259, 261, 262, 271, 272, 274, 279, 287, 293, 294, 295, 299, 301, 302, 306, 307, 308, 309, 315, 316, 323, 324, 325, 326, 327, 332, 333, 334, 335, 337, 338, 339, 342, 343, 344, 345, 346, 348, 350, 351, 353, 356, 363, 365, 368, 371, 375, 379, 381 and 244311-250683.
- 31 Staphylococcus aureus subsp. aureus N315 2, 5, 7, 8, 9, 10, 13, 16, 19, 22, 25, 27, 31, 32, 33, 35, 36, 38, 39, 40, 41, 45, 46, 47, 48, 50, 51, 52, 55, 62, 63, 67, 71, 73, 81, 83, 84, 85, 90, 91, 92, 93, 95, 98, 100, 101, 105, 106, 111, 113, 117, 119, 120, 124, 131, 133, 134, 138, 139, 143, 146, 147, 149, 152, 153, 156, 160, 161, 162, 166, 169, 171, 172, 174, 177, 179, 180, 181, 190, 192, 203, 204, 205, 207, 208, 213, 214, 217, 218, 222, 226, 228, 231, 232, 236, 238, 240, 242, 244, 245, 247, 248, 252, 254, 256, 259, 260, 261, 262, 270, 271, 272, 274, 275, 279, 287, 293, 294, 299, 301, 302, 305, 306, 307, 308, 309, 311, 316, 317, 323, 324, 325, 326, 327, 332, 333, 334, 335, 337, 339, 340, 342, 343, 344, 345, 346, 348, 349, 351, 353, 354, 356, 363, 365, 368, 371, 375, 379, 381 and 250684-257140.
- 32 Streptococcus Pneumoniae R6 2, 3, 5, 6, 10, 13, 14, 17, 20, 21, 22, 23, 25, 26, 27, 30, 31, 32, 33, 35, 36, 37, 38, 39, 40, 41, 46, 47, 48, 49, 50, 52, 55, 56, 62, 63, 67, 73, 77, 81, 83, 84, 85, 87, 90, 91, 92, 94, 95, 100, 101, 102, 105, 106, 111, 112, 114, 115, 116, 117, 119, 123, 124, 126, 133, 136, 138, 143, 145, 146, 147, 149, 152, 156, 160, 161, 164, 166, 168, 169, 171, 172, 174, 175, 176, 177, 179, 180, 190, 192, 203, 204, 205, 208, 209, 213, 214, 217, 218, 223, 226, 228, 229, 232, 233, 235, 236, 238, 239, 242, 244, 245, 246, 247, 248, 249, 252, 255, 256, 257, 258, 259, 260, 261, 262, 264, 268, 271, 272, 274, 279, 282, 283, 284, 287, 295, 296, 297, 298, 299, 300, 302, 303, 305, 306, 307, 309, 311, 312, 314, 315, 316, 320, 321, 323, 324, 325, 326, 327, 329, 333, 335, 338, 340, 341, 344, 345, 348, 350, 351, 352, 353, 356, 357, 359, 365, 368, 371, 372, 373, 375, 377, 379, 380, 382, 384, 385 and 257141-265301.
- 33 Streptococcus pneumoniae TIGR4 2, 10, 13, 25, 27, 33, 46, 48, 50, 52, 55, 62, 63, 67, 73, 81, 84, 91, 101, 105, 106, 111, 119, 149, 152, 160, 161, 176, 177, 164, 166, 168, 169, 171, 172, 175, 179, 180, 190, 205, 208, 213, 214, 218, 228, 236, 242, 244, 246, 262, 268, 271, 272, 274, 297, 299, 306, 321, 323, 324, 325, 327, 329, 333, 340, 345, 348, 351, 353, 356, 359, 365, 368, 371, 372, 375, 380 and 265302-266788.
- 34 Streptococcus pyogenes M1 GAS 3, 5, 8, 10, 21, 22, 25, 27, 32, 37, 38, 39, 40, 43, 49, 90, 95, 96, 106, 116, 126, 129, 138, 163, 164, 168, 175, 261, 262, 176, 180, 226, 232, 244, 246, 259, 268, 283, 295, 296, 297, 299, 306, 309, 316, 321, 329, 330, 333, 348, 349, 359, 372, 379, 380 and 266789-269521.
- 35 Streptococcus pyogenes MGAS315 3, 8, 10, 13, 20, 22, 25, 27, 31, 32, 33, 37, 38, 40, 46, 48, 52, 55, 62, 67, 73, 84, 90, 91, 105, 106, 113, 116, 175, 176, 129, 138, 152, 160, 164, 166, 168, 177, 179, 180, 186, 190, 192, 205, 208, 211, 213, 214, 218, 226, 229, 232, 236, 242, 244, 246, 262, 268, 271, 272, 274, 282, 283, 295, 296, 297, 299, 306, 309, 312, 321, 323, 324, 325, 327, 329, 333, 340, 345, 348, 349, 353, 356, 359, 372, 379, 380, 381 and 269522-272357.
- 36 Streptococcus pyogenes MGAS8232 3, 4, 8, 10, 13, 21, 22, 25, 27, 31, 33, 37, 38, 39, 40, 46, 48, 52, 55, 62, 67, 73, 84, 90, 91, 95, 105, 106, 113, 116, 129, 138, 168, 152, 160, 163, 164, 166, 175, 176, 177, 179, 180, 190, 205, 208, 213, 214, 218, 226, 232, 236, 242, 244, 246, 247, 259, 260, 261, 262, 268, 271, 272, 274, 295, 296, 297, 299, 306, 307, 309, 316, 321, 323, 324, 325, 327, 329, 330, 333, 337, 340, 344, 345, 348, 349, 353, 356, 359, 363, 372, 379, 380, 381 and 272358-275553.
- 37 Streptococcus 10, 13, 25, 27, 31, 33, 46, 48, 52, 55, 62, 67, 73, 84, 91,

- pyogen 164, 166, 168, 175, 176, 177, 179, 180, 190, 205, 208, 213, 214,
es SSI-1 242, 218, 236, 244, 246, 262, 268, 271, 272, 274, 297, 299, 306,
324, 325, 327, 329, 321, 323, 105, 113, 152, 160, 333, 340, 345,
348, 353, 356, 359, 372, 380, 381 and 275554-276703.
- 38 *Treponema s* 3, 10, 13, 48, 52, 57, 59, 67, 81, 84, 86, 90, 91, 121, 131, 134,
ubsp. 174, 175, 176, 184, 218, 228, 231, 235, 236, 243, 261, 262, 269, 272, 306,
pallidum str. 289, 291, 295, 299, 312, 324, 329, 332, 333, 340
Nichols , 345, 356, 358 and 276704-277654.
- 39 *Yersinia* 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 12, 13, 16, 18, 19, 21, 22,
pestis 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 36, 37, 39, 40, 41, 54,
42, 43, 45, 46, 47, 48, 51, 52, 53, 55, 57, 58, 61, 62, 63, 67,
68, 70, 71, 73, 75, 76, 78, 82, 84, 85, 87, 88, 89,
90, 91, 93, 94, 95, 98, 99, 101, 102, 103, 105, 106, 107,
108, 111, 112, 113, 114, 115, 116, 117, 120, 121, 122, 123, 124,
125, 126, 129, 130, 131, 132, 133, 134, 135, 136, 138, 140, 141,
142, 143, 146, 148, 149, 151, 152, 153, 154, 155, 156, 160, 164,
165, 166, 167, 169, 171, 172, 174, 175, 176, 177, 178, 179, 180,
182, 184, 186, 187, 188, 190, 191, 192, 193, 196, 197, 198, 199,
200, 201, 202, 203, 205, 206, 208, 209, 211, 213, 214, 215, 217,
218, 219, 220, 221, 222, 224, 225, 226, 227, 229, 230, 232, 233,
234, 236, 237, 238, 239, 240, 241, 242, 243, 244, 245, 250, 251,
252, 253, 255, 256, 257, 258, 259, 260, 262, 263, 264, 270,
271, 272, 274, 276, 279, 280, 281, 282, 283, 286, 287, 289, 291,
292, 293, 295, 296, 298, 299, 300, 301, 302, 304, 306, 307, 308,
309, 311, 314, 315, 317, 319, 321, 322, 323, 324, 325, 326, 327,
329, 330, 331, 333, 334, 335, 336, 337, 340, 341, 342, 343, 344, 345,
346, 348, 349, 350, 351, 352, 353, 354, 355, 356, 357,
358, 359, 363, 364, 365, 367, 368, 370, 372, 373, 374, 376, 377, 378,
379, 380, 381, 382, 383, 384 and 277655-287825.
- 40 *Yersinia* 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 12, 13, 16, 18, 19, 20,
pestis 21, 22, 25, 26, 27, 28, 29, 31, 32, 33, 34, 36, 37, 39, 40, 41, 53, 54,
KIM 42, 43, 45, 46, 47, 48, 51, 52, 55, 57, 58, 61, 62, 63, 65, 67, 68, 70,
71, 72, 73, 75, 76, 78, 84, 85, 87, 88, 89, 90, 91, 93, 94, 95, 97, 99,
101, 102, 103, 105, 106, 107, 108, 111, 112, 113, 114, 115, 117, 118,
120, 121, 122, 123, 124, 125, 126, 129, 130, 131, 132, 133, 134, 135,
136, 138, 140, 142, 143, 146, 147, 148, 149, 151, 152, 153, 154,
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